Supporting Information

Contrib	uting centers	Provided case data (N)	Prospective included cases (N)	Retrospective included cases (N)	Excluded cases (N) 12	
All cent	ers	209	175	22		
Austral	ia MotherSafe, Sidney	10	8	1	1 ^a	
France	French Network of Pharmacovigilance Centers, Terappel	99 35	78 32	14 3	7 0	
	Centre de Référence sur les Agents Tératogènes, Paris	64	46	11	7 ⁶	
Germar	y Institute for Clinical Teratology and Drug Risk Assessment in Pregnancy, Berlin	29	24	5	0	
Israel	The Israeli Teratology Information Service, Jerusalem	16	15	1	0	
Italy	Centro di Riferimento Regionale di Tossicologia Perinatale, Firenze	4 1	4 1	0 0	0 0	
	Teratology Information Service, Telefono Rosso, Catholic University of Sacred Heart, Rome	2	2	0	0	
	TIS PADUA-CEPIG, Clinical Genetics Unit, Department of Women's and Children's Health, University of Padua	1	1	0	0	
Japan	Japan Drug Information Institute in Pregnancy (JDIIP), Tokyo	5	5	0	0	
Netherl	ands Teratology Information Service of the Netherlands Pharmacovigilance Centre Lareb	2	1	0	1 [°]	
Spain	Clinical Pharmacology Service – Fundació Institut Català de Farmacologia, TIS Barcelona	10 1	10 1	0 0	0 0	

Table S1_SuppInfo. Contributing centers and provided data.

	Servicio de Información Telefónica sobre Teratógenos Español (SITTE) – Service for health professionals, Madrid	9	9	0	0
Switzer	land Swiss Teratogen Information Service, Division de Pharmacologie Clinique, Centre hospitalier universitaire vaudois, Lausanne	5	5	0	0
Turkey	Medical Pharmacology Outpatient Clinics, Marmara University Education and Research Hospital, Istanbul	11 3	10 3	0 0	1 0
	Terafar – Izmir Katip Celebi University Teratology Information, Training and Research Center, Izmir	4	3	0	1 ^c
	Trabzon-TIS — Teratology Information Service, Department of Pharmacology, Karadeniz Technical University, Trabzon	4	4	0	0
United	Kingdom UK Teratology Information Service, Newcastle upon Tyne	13	12	1 (twin pregnancy)	0
USA	MotherToBaby, Center for Better Beginnings, University of California San Diego	5	3 (including one twin pregnancy)	0	2 ^ª

^a Exclusion because estimated date of birth was after August 1st, 2019. ^b Exclusion of four cases because crucial data were missing, exclusion of three cases as duplicates to other French cases. ^c Exclusion because of preconceptional exposure only. ^d Exclusion because of preconceptional exposure only in one case, exclusion of another because crucial date were missing.

Table S2_SuppInfo. Reference charts for neonatal growth parameters.

Country and center	Type of growth chart and references ¹		
Australia MotherSafe, Sidney	International tool: (Villar et al., 2014) and (Villar et al., 2015), underlying data can be accessed via https://intergrowth21.tghn.org/standards-tools/		
France French Network of Pharmacovigilance Centers, Terappel	Country-specific tool: https://www.audipog.net/Courbes-morpho , brief summary of methods on the website and in (Vendittelli, Rivière, Pinguier, & Claris, 2008)		
Centre de Référence sur les Agents Tératogènes, Paris			
Germany Institute for Clinical Teratology and Drug Risk Assessment in Pregnancy, Berlin	Publication: (M. Voigt et al., 2014) ²		
Israel The Israeli Teratology Information Service, Jerusalem	Publication: (Dollberg, Haklai, Mimouni, Gorfein, & Gordon, 2005) ²		
Italy Centro di Riferimento Regionale di Tossicologia Perinatale, Firenze	Country-specific tool: (Bertino et al., 2010), underlying data can be accessed via		
TIS PADUA-CEPIG, Clinical Genetics Unit, Deparment of Women's and Children's Health, University of Padua	http://www.inescharts.com/		
Teratology Information Service, Telefono Rosso, Catholic University of Sacred Heart, Rome			
Japan Japan Drug Information Institute in Pregnancy (JDIIP), Tokyo	Country-specific tool: (Itabashi et al., 2010), underlying data can be accessed via http://jspe.umin.jp/medical/keisan.html		
Netherlands Teratology Information Service of the Netherlands Pharmacovigilance Centre Lareb	Publication: (Hoftiezer et al., 2019) ² , underlying data can be accessed via <u>https://www.perined.nl/geboortegewichtcurven</u>		
Spain Clinical Pharmacology Service – Fundació Institut Català de Farmacologia, TIS Barcelona	Publication: (Generalitat de Catalunya Departament de Salut, 2008) ^{2,4} , accessible via https://scientiasalut.gencat.cat/bitstream/handle/11351/1209/corbes_referencia_pes_londgitu		
Servicio de Información Telefónica sobre Teratógenos Español (SITTE) – Service for health professionals, Madrid	<u>d nounats catalunya_2010.pdf?sequence=1</u>		
Switzerland Swiss Teratogen Information Service, Division de Pharmacologie Clinique, Centre hospitalier universitaire vaudois, Lausanne	Publication: (M Voigt et al., 2006) ⁵ , see also <u>https://cdn.paediatrieschweiz.ch/production/uploads/2020/05/Perzentilen_2012_09_15_SGP_d-1.pdf</u>		

Turkey Medical Pharmacology Outpatient Clinics, Marmara University Education and Research Hospital, Istanbul	Publication: (Kurtoğlu et al., 2012)
Terafar – Izmir Katip Celebi University Teratology Information, Training and Research Center, Izmir	
Trabzon-TIS — Teratology Information Service, Department of Pharmacology, Karadeniz Technical University, Trabzon	
United Kingdom UK Teratology Information Service, Newcastle upon Tyne	Publication: (Cole, Williams, & Wright, 2011)
USA MotherToBaby, Center for Better Beginnings, University of California San Diego	International tool: (Villar et al., 2014) and (Villar et al., 2015), the underlying data can be accessed via https://intergrowth21.tghn.org/standards-tools

¹Country-specific reference standards provided SDS either directly or indirectly by a precise quantile (France, Italy, Japan), or they provided the parameters (or data to approximate the parameters) for applying the LMS method by Cole (Cole & Green, 1992), as in the reference standards for Germany, Israel, Netherlands, Spain, Switzerland, Turkey and UK. The LMS method is a transformation from skewed data to standard normal, using the three parameters skewness (L), median (M) and coefficient of variation (S). Last date of access for web-based tools or data: 28 April 2023. ² Without LMS parameters, but including data to approximate the parameters for the LMS method. ³ Data accessed on 8 June 2022; version 1.1 was used for study cases. ⁴ Growth parameters were collected in Catalonia only. In this study, they were used for all Spanish study cases. ⁵Statistical methodology uses normal distribution.

References regarding neonatal growth charts in Table S2_SuppInfo

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Table S3 SuppInfo. SDS means for birth weight (BW) and neonatal head circumference (HC) based on international vs. country-specific reference charts.

SDS means for BW and neonatal HC based on international (Intergrowth-21st) vs. country-specific reference charts show large differences. In Japanese newborns, for example, the difference of SDS means for BW amounts to +0.74 SDS, the difference of SDS means for HC to +0.42 SDS. In other words, SDS means of Japanese growth parameters in the cohort analyzed here are much higher if based on country-specific reference standards than if based on Intergrowth-21st standards. For infants from most other countries the opposite is true: SDS means are considerably smaller if based on country-specific reference standards. For example, the SDS mean for BW in German newborns is -0.34 SDS smaller according to country-specific reference charts than according to Intergrowth-21st, and the SDS mean for HC decreases by -0.9 SDS.

		Birth we	eight: SDS ¹ mean		Neonatal head circumference: SDS ¹ mean			
COUNTRY	Cases (N)	International reference charts	Country-specific reference charts	Difference	Cases (N)	International reference charts	Country-specific reference charts	Difference
Australia	4	+0.46	N/A	N/A	4	+0.08	N/A	N/A
France	64	-0.12	-0.22	-0.10	31	+0.24	-0.21	-0.45
Germany	20	-0.16	-0.50	-0.34	19	+0.41	-0.49	-0.90
Israel	14	-0.35	-0.26	+0.09	0	N/A	N/A	N/A
Italy	2	-1.22	-1.49	-0.27	1	+0.44	+0.28	-0.16
Japan	5	-0.06	+0.69	+0.74	5	+0.12	+0.54	+0.42
Netherlands	1	-0.06	-0.65	-0.58	0	N/A	N/A	N/A
Spain	10	-0.55	-0.56	-0.01	4	-0.18	-0.45	-0.28
Switzerland	5	+0.13	-0.25	-0.37	3	+1.48	+0.44	-1.04
Turkey	6	+0.85	+0.94	0.09	1	+3.04	+2.44	-0.60
UK	9	-0.29	-0.37	-0.07	2	-0.06	-0.32	-0.26
USA	4	-0.26 ²	N/A	N/A	3	+0.91 ²	N/A	N/A

¹ SDS: Standard deviation score. ² Since reference charts for twins were not available, percentiles for one set of twins were calculated according to singleton reference charts. If they are excluded from the analysis, SDS mean for BW (N=2) is -0.83 and for neonatal HC (N=1) -0.18.

Table S4_SuppInfo. Overview of prospectively ascertained study cases included for defined outcomes.

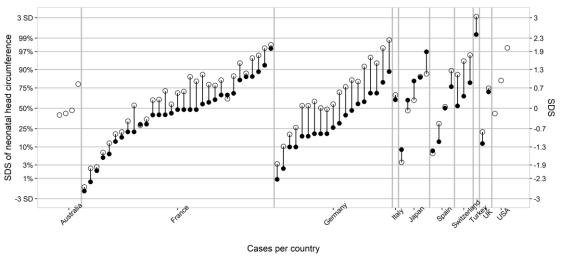
	Study inclusion criteria	N
All prospectively ascertained	Any exposure to modafinil between 2+0 weeks after last menstrual period and delivery.	175 pregnancies
cases	Estimated date of birth (EDOB) before August 2019.	
	Data on pregnancy outcome and congenital malformations available.	
Outcome analysis	Additional criteria for inclusion in specific analysis	Ν
Cumulative incidences of	Start of modafinil exposure before study entry.	174 pregnancy outcomes
pregnancy outcome		
Major congenital malformations	First trimester exposure to modafinil.	150 pregnancy outcomes
(MCA)	No diagnosed chromosomal/genetic disorder of the fetus/newborn.	
	Fetus affected with MCA or live birth.	
Percentile values and standard	Live birth.	144 newborns
deviation scores (SDS) for birth	Measurement of BW available.	
weight (BW)	Data on gestational age at birth and neonatal sex available.	
Percentile values and standard	Live birth.	73 newborns
deviation scores (SDS) neonatal	Measurement of neonatal HC available.	
head circumference (HC)	Data on gestational age at birth and neonatal sex available.	
Potential dose-dependent effects	Live birth.	121 newborns
on BW	Measurement of BW available.	
	Data on cumulative modafinil dosage, gestational age at birth and neonatal sex available.	
Potential dose-dependent effects	Live birth.	59 newborns
on HC	Measurement of neonatal HC available.	
	Data on cumulative modafinil dosage, gestational age at birth and neonatal sex available.	
Postnatal Growth	Live birth.	4 infants
	Modafinil exposure in all three trimesters.	
	At least one measurement of weight and HC after the first month of life.	
	Data on gestational age at birth and infant sex available.	

Pregnancy outcome, completed GW ¹ at delivery, sex	Modafinil exposure (GW ¹ ; dosage)	Indication for modafinil treatment	Major congenital anomalies (MCA) ² , additional minor anomalies and/or remarks
Live birth, GW 41, male	GW 0 – 5+4; 400 mg/d	Idiopathic hypersomnia	Unilateral foot malformation: cutaneous syndactyly (4th-5th toes) and supernumerary metatarsal bone. Mild facial dysmorphism, jaw-winking. Epidermoid cyst in skull bone. Orbital cyst. Bilateral inguinal hernia. Relevant vesicoureteral reflux. Genetic testing provided unremarkable results.
Live birth, GW 41, female	GW 0 – 39+3; during 1 st trimester 200 mg/d	Narcolepsy	Pierre Robin sequence with retrognathia, glossoptosis and cleft palate. Metatarsus varus.
Live birth, GW 38, female	GW 0 – 2+3; 300 mg/d	Narcolepsy	Esophageal atresia. Neonatal death within the first week of life.

Table S5_SuppInfo. Infants with major congenital anomalies (MCA).

¹ GW: gestational weeks. ² None of the affected newborns was exposed to concurrent medication with known teratogenic effects. Study cases with chromosomal aberrations are not included.





O International reference chart (Intergrowth-21st)

Country-specific reference chart

Figure S1_SuppInfo. Standard deviation scores (SDS) and percentiles (in %) of individual neonatal head circumference (HC) in different countries according to international (Intergrowth-21st) vs. country-specific reference charts. No severe microcephaly (head circumference less than -3 SD) was detected, regardless of whether country-specific or international percentiles were used.



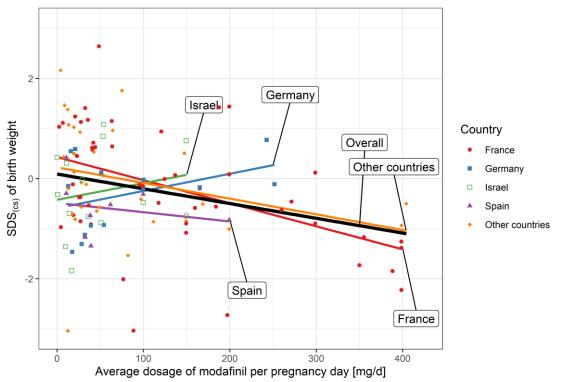


Figure S2 Supplnfo. Potential dose-dependent effects of modafinil on birth weight (BW): Countryspecific and overall correlations in a linear regression model. For statistical details of this model, see Table S6_SuppInfo (presented below Figure S3_SuppInfo). The **black line** represents the regression line for all cases combined (N=121). Other countries refers to all countries with less than ten observations for BW and country-specific reference charts, i.e. Italy, Japan, Netherlands, Switzerland, Turkey and United Kingdom. SDS(cs): Standard deviation scores based on country-specific reference charts taking into account neonatal sex and GW at delivery. If an average dosage per pregnancy day of zero (i.e. no modafinil exposure) is assumed, the model obtains an overall SD score of 0.054 for BW (95% CI, -0.2 to 0.3). For all countries combined, a decreasing slope is obtained, also for France, Spain and the group of countries contributing less than ten cases. In contrast, the slope is increasing for Germany and Israel. If the findings for Germany are compared to France, the slope increases by the factor 0.77 SDS (France -0.43, Germany +0.34, 95% CI, 0.07 to 1.48) with each 100 mg increase of average dosage per pregnancy day. However, the main reason for this effect are the SDS differences in cases with low dosages per pregnancy day: In these cases, the SDS of BW are considerably lower in Germany than in France, resulting in a considerably lower intercept. In fact, the intercept of Germany (i.e. the SDS for BW with an assumed modafinil exposure of zero) is lowered by -0.91 SDS (France: 0.32, Germany: -0.61, 95% CI, -1.77 to -0.05). The data for Israel are similar, but results lack statistical significance due to low sample size. Furthermore, Germany (as well as Israel) contributed no cases with very high (\geq 300 mg) average dosages per pregnancy day. In an additional model, we eliminated the impact of sample-specific intercept differences between countries by using the same intercept for each country (see Figure S3_SuppInfo).



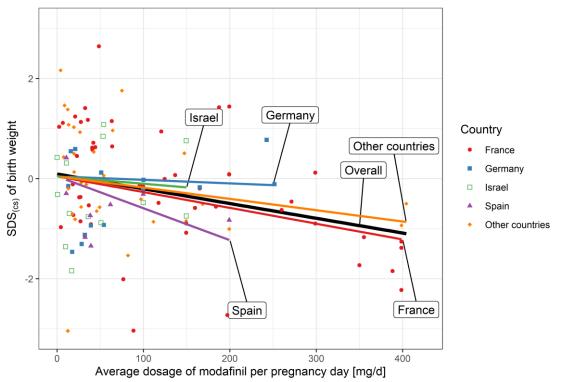


Figure S3_SuppInfo. Potential dose-dependent effects of modafinil on birth weight (BW): Additional linear regression model to eliminate the impact of sample-specific intercept differences between countries. For each country, the same intercept is used, i.e. 0,043 SDS. The **black line** represents the regression line for all cases combined (N=121). In this model, the slopes of Germany and Israel no longer increase and the differences between France and Germany are no longer significant. For statistical details of this model, see Table S6_SuppInfo (presented below this Figure). Other countries refers to all countries with less than ten observations for BW and country-specific reference charts, i.e. Italy, Japan, Netherlands, Switzerland, Turkey and United Kingdom. SDS(cs): Standard deviation scores based on country-specific reference charts taking into account neonatal sex and GW at delivery.

Table S6_SuppInfo. Coefficient tables and summary statistics for the linear models of potential dose-dependent effects on birth weight (for Figures S2_SuppInfo and S3_SuppInfo).

Note that avg_dosage refers to cumulative modafinil dosage in mg divided by the duration of pregnancy in days. Thus, the effects reported in the tables above refer to a 1 mg increase in dosage, while in the main text 100 mg effects were reported for better readability. For separations by country, the reference is always France. All confidence intervals have a level of 95%.

Model for birth weight: all countries combined (Figure S2_SuppInfo)

Coefficient tables and summary statistics for all cases combined (N=121), represented in Figure S2_SuppInfo by the black regression line.

term	estimate	std.error	statistic	p.value	conf.low	conf.high
(Intercept)	0.0544	0.126	0.432	0.667	-0.195	0.304
avg_dosage	-0.00275	0.000864	-3.19	0.00184	-0.00446	-0.00104

	summary statistics
r.squared	0.0786
adj.r.squared	0.0709
sigma	1.01
statistic	10.2
p.value	0.00184
df	1
logLik	-172
AIC	350
BIC	359
deviance	122
df.residual	119
nobs	121

Model for birth weight: full interaction by country (slope and intercept, Figure S2_SuppInfo) Coefficient tables and summary statistics for all countries separately, represented in Figure S2_SuppInfo by the colored regression lines.

term	estimate	std.error	statistic	p.value	conf.low	conf.high
(Intercept)	0.324	0.211	1.54	0.127	-0.0936	0.741
avg_dosage	-0.0043	0.00117	-3.67	0.000373	-0.00662	-0.00198
country Germany	-0.913	0.434	-2.11	0.0375	-1.77	-0.0536
country Israel	-0.746	0.431	-1.73	0.0867	-1.6	0.109
country other	-0.111	0.308	-0.36	0.72	-0.721	0.499
country Spain	-0.808	0.556	-1.46	0.148	-1.91	0.292
avg_dosage:country Germany	0.00774	0.00357	2.16	0.0326	0.000654	0.0148
avg_dosage:country Israel	0.00763	0.00562	1.36	0.178	-0.00351	0.0188
avg_dosage:country other	0.00158	0.00203	0.78	0.437	-0.00243	0.0056
avg_dosage:country Spain	0.00245	0.00617	0.398	0.692	-0.00977	0.0147

	summary statistics
r.squared	0.155
adj.r.squared	0.0863
sigma	1
statistic	2.26
p.value	0.0232
df	9
logLik	-167
AIC	356
BIC	387
deviance	112
df.residual	111
nobs	121

Model for birth weight: interaction by country on slope, but not on intercept (Figure S3_SuppInfo)

Coefficient tables and summary statistics for the regression lines in Figure S3_SuppInfo. In the model with full interaction (Figure S2_SuppInfo), the country-specific intercepts, i.e. the estimated SDS means for BW at a modafinil dosage of 0 mg, differ by almost 1 SD. This difference is quite large and therefore likely to be an overestimate, having a substantial effect on the estimated slopes. While no country-specific differences in intercept are also unlikely, this second model allows the reader to consider the spectrum between these two extremes.

term	estimate	std.error	statistic	p.value	conf.low	conf.high
(Intercept)	0.0434	0.131	0.331	0.741	-0.216	0.303
avg_dosage	-0.00314	0.000962	-3.26	0.00145	-0.00505	-0.00123
avg_dosage:country Germany	0.00246	0.00249	0.986	0.326	-0.00248	0.0074
avg_dosage:country Israel	0.00169	0.00413	0.41	0.683	-0.00649	0.00988
avg_dosage:country other	0.0012	0.00153	0.787	0.433	-0.00183	0.00424
avg_dosage:country Spain	-0.00321	0.00436	-0.737	0.463	-0.0118	0.00542

summary statistics
0.0961
0.0568
1.02
2.44
0.0381
5
-171
356
376
120
115
121

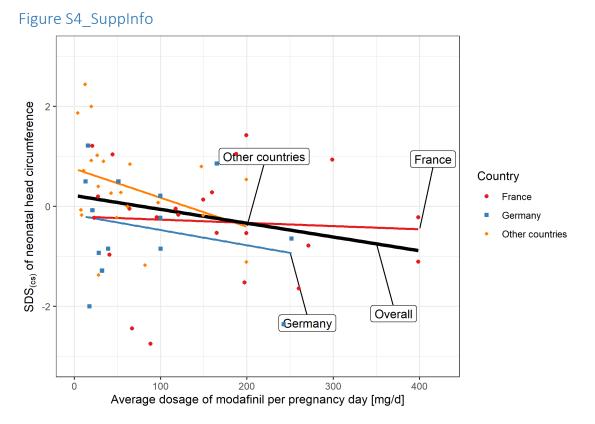


Figure S4_SuppInfo. Potential dose-dependent effects of modafinil on neonatal head circumference (HC): Country-specific and overall correlations in a linear regression model. For statistical details of this model, see Table S7_SuppInfo (presented below Figure S5_SuppInfo). The **black line** represents the regression line for all cases combined (N=59). Other countries refers to all countries with less than ten observations for HC, i.e. Italy, Japan, Spain, Switzerland, Turkey and United Kingdom. SDS(cs): Standard deviation scores based on country-specific reference charts and taking into account neonatal sex and GW at delivery. The linear model to evaluate interactions between intercept and country as well as slope and country finds only small, non-significant differences in the intercept and slope of the different countries. Taking into account all countries and assuming an average dosage per pregnancy day of zero (i.e. no modafinil exposure), our linear model obtained an SDS of 0.22 (95%CI, -0.19 to 0.62). In an additional model, we eliminated the impact of sample-specific intercept differences between countries by using the same intercept for each country (see Figure S5_SuppInfo).

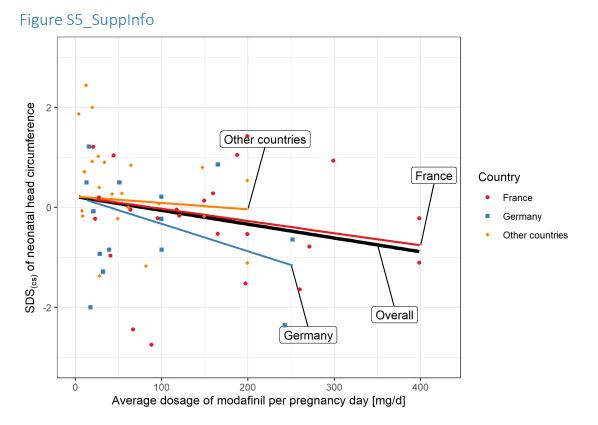


Figure S5_SuppInfo: Potential dose-dependent effects of modafinil on neonatal head circumference (HC). Additional linear regression model to eliminate the impact of sample-specific intercept differences between countries. The black line represents the regression line for all cases combined (N=59). Using a common intercept for all countries results in even smaller differences in the slope of the different countries than the original model (Figure S4_SuppInfo). For statistical details of this model, see Table S7_SuppInfo (presented below this Figure). Other countries refers to all countries with less than ten observations for HC, i.e. Italy, Japan, Spain, Switzerland, Turkey and United Kingdom. SDS(cs): Standard deviation scores based on country-specific reference charts and taking into account neonatal sex and GW at delivery.

Table S7_SuppInfo. Coefficient tables and summary statistics for the exploratory linear models of potential dose-dependent effects on neonatal head circumference (for Figures S4_SuppInfo and S5_SuppInfo).

Note that avg_dosage refers to cumulative modafinil dosage in mg divided by the duration of pregnancy in days. Thus, the effects reported in the tables refer to a 1 mg increase in dosage, while in the main text 100 mg effects were reported for better readability. For separations by country, the reference is always France. All confidence intervals have a level of 95%.

Model for neonatal HC: all countries combined (Figure S4_SuppInfo)

Coefficient tables and summary statistics for all cases combined (N=59), represented in Figure S4_SuppInfo by the black regression line.

term	estimate	std.error	statistic	p.value	conf.low	conf.high
(Intercept)	0.217	0.203	1.07	0.29	-0.19	0.623
avg_dosage	-0.00276	0.00143	-1.92	0.0595	-0.00563	0.000114

	summary statistics
r.squared	0.0609
adj.r.squared	0.0444
sigma	1.06
statistic	3.7
p.value	0.0595
df	1
logLik	-86.2
AIC	178
BIC	185
deviance	64.2
df.residual	57
nobs	59

Model for neonatal head circumference: full interaction by country (slope and intercept, Figure S4 SuppInfo)

Coefficient tables and summary statistics for all countries separately, represented in Figure S4_SuppInfo by the colored regression lines.

term	estimate	std.error	statistic	p.value	conf.low	conf.high
(Intercept)	-0.201	0.378	-0.531	0.598	-0.958	0.557
avg_dosage	-0.000643	0.00198	-0.325	0.747	-0.00462	0.00333
country Germany	0.0327	0.554	0.059	0.953	-1.08	1.14
country other	0.954	0.494	1.93	0.0587	-0.0364	1.94
avg_dosage:country Germany	-0.0024	0.00403	-0.595	0.554	-0.0105	0.00568
avg_dosage:country other	-0.00515	0.00423	-1.22	0.229	-0.0136	0.00333

	summary statistics
r.squared	0.166
adj.r.squared	0.0869
sigma	1.04

	summary statistics
statistic	2.1
p.value	0.0792
df	5
logLik	-82.7
AIC	179
BIC	194
deviance	57.1
df.residual	53
nobs	59

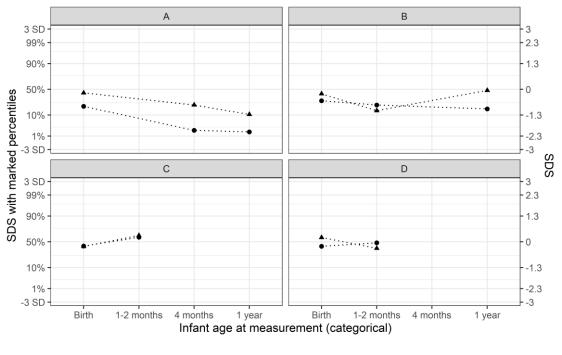
Model for neonatal HC: interaction by country on slope, but not on intercept (Figure S5_SuppInfo)

Coefficient tables and summary statistics for the regression lines in Figure S5_SuppInfo. In the model with full interaction (Figure S4_SuppInfo), the country-specific intercepts, i.e. the estimated SDS means for neonatal HC at a modafinil dosage of 0 mg, differ by almost 1 SD. This difference is quite large and therefore likely to be an overestimate, having a substantial effect on the estimated slopes. While no country-specific differences in intercept are also unlikely, this second model allows the reader to consider the spectrum between these two extremes.

term	estimate	std.error	statistic	p.value	conf.low	conf.high
(Intercept)	0.218	0.214	1.02	0.313	-0.211	0.647
avg_dosage	-0.00244	0.00148	-1.65	0.106	-0.00542	0.000533
avg_dosage:country Germany	-0.00303	0.00276	-1.1	0.277	-0.00857	0.0025
avg_dosage:country other	0.00117	0.00304	0.384	0.703	-0.00493	0.00727

	summary statistics
r.squared	0.0874
adj.r.squared	0.0376
sigma	1.07
statistic	1.76
p.value	0.166
df	3
logLik	-85.4
AIC	181
BIC	191
deviance	62.4
df.residual	55
nobs	59

Figure S6 SuppInfo



▲ Head circumference in SDS ● Infant weight in SDS

Figure S6_SuppInfo. Postnatal growth curves for weight and head circumference (HC). Postnatal growth of four children from the United States [A], Australia [B] and Germany [C and D], none of them with MCA. Weight and HC in SDS (standard deviation scores) with marked percentiles, infant age in categories. For child A, percentile ranges of infant weight and HC were continuously decreasing with infant age: Over the course of the first year of life, the percentile for weight decreased from the 20th to below the 2nd percentile, and for HC from just below median to the 10th percentile. Additionally, developmental delay was reported. For the other three children, no overall trend of decreasing percentile ranges was identified.