

EURAP

An International Antiepileptic Drugs and Pregnancy Registry

Interim Report – MAY 2023

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BACKGROUND

A number of independent groups with experience and interest in maternal and foetal well-being in association with maternal use of antiepileptic drugs (AEDs)* have agreed on a prospective international multi-centre study of pregnancies with AEDs. Data from all participating groups are shared in a Central Registry of Antiepileptic Drugs and Pregnancy (EURAP). EURAP was established in the first centres in some European countries and has since then gradually expanded to include more centres and countries now involving also Asia, Oceania, Latin America and Africa. The EURAP Study protocol has been updated in June 2021 and can be found on www.eurapinternational.org

* since integrated in the project name and acronym we maintain in this document the term AED rather than the now proposed term antiseizure medication, ASM.

OBJECTIVE OF EURAP

The primary objective of EURAP is to evaluate and determine the comparative risk of major foetal malformations following intake of AEDs and their combinations during pregnancy.

METHODS

EURAP is an observational study. Women taking AEDs at the time of conception, irrespective of the indication, may be included. To avoid selection bias, only pregnancies recorded before foetal outcome is known and within week 16 of gestation are included in the prospective risk assessment. Cases ascertained later in pregnancy are recorded as retrospective cases, as they may provide signals, but are not included in the comparative risk evaluation.

Information on patient's demographics, type of epilepsy, seizure frequency, family history of malformations, drug therapy and of other potential risk factors is obtained, and follow-up data are collected once at each trimester, at birth and at one year after delivery.

Networks of reporting physicians have been established in countries taking part in the collaboration. During the course of the pregnancy, and the follow-up time after delivery, the participating physician enters data into five Subforms (Subforms A-E) for each patient.

Subform A is completed on enrolment of the patient, Subform B after the first trimester, Subform C after the second trimester, Subform D within three months after delivery, and Subform E within 14 months after birth. Immediately after completion, each Subform is submitted to the national coordinator for review. The national coordinator transfers the reviewed and accepted Subform to the Central EURAP Registry in Milan, Italy.



EVALUATION OF OUTCOME

The physician records descriptively abnormalities observed in the offspring. The final assessment and classification of the type of malformation is the responsibility of the Central Project Commission (CPC). In order to facilitate a uniform and objective assessment, reports of malformations are assessed regularly by an outcome assessment committee, which is kept blinded with respect to the type of exposure.

INTERIM REPORT

EURAP was implemented in the first two countries in Europe in 1999 and has since then grown to include countries from Europe, Oceania, Asia, Latin America and Africa. This development is reflected by increasing numbers of enrolled pregnancies. The development since 1999 is illustrated in Figure 1.

<u>Figure 1</u>. Number of Participating Countries and Pregnancies Reported to the Central Registry by March, 2023.





The present report is based on data available in the Central Registry by MAY 25th, 2023.

At that time more than 1,500 reporting physicians from 47 countries had contributed cases to the Central Registry. Countries that have contributed at least 10 pregnancies in the current report are listed in Table 1.

Table 1. Countries that have contributed at least 10 pregnancies in the current report (n=38).

COUNTRY	National Coordinator (or referring physician*)	Date of joining the Registry		
Argentina	Silvia Kochen	2002		
Australia	Frank Vajda	2000		
Austria	Gerhard Luef	2000		
Belarus	Halina Navumava*	2008		
Belgium	Dick Lindhout & Eugène van Puijenbroek	2002		
Chile	Alejandro De Marinis	2002		
China	Weiping Liao	2006		
Croatia	Dinko Vitezic	2002		
Czech Republic	Jana Zarubova	2001		
Denmark	Anne Sabers	2000		
El Salvador	Ovidio Solano Cabrera*	2017		
Estonia	Aleksei Rakitin*	2019		
Finland	Reetta Kälviäinen	2003		
France	Marion Quirins*	2000		
Georgia	Sofia Kasradze; Nino Gogatishvili*	2000		
Germany	Bettina Schmitz	2000		
Hong-kong	Patrick Kwan	2002		
India	Sanjeev Thomas	2001		
Iran	Nasim Tabrizi	2018		
Israel	Lilach Goldstein	2000		
Italy	Barbara Mostacci	2000		
Japan	Hideyuki Ohtani	2001		
Lithuania	Ruta Mameniskiene	2002		
Macedonia	Gordana Kiteva Trencevska	2001		
Netherlands	Dick Lindhout & Eugène van Puijenbroek	2002		
Norway	Silje Alvestad	2000		
Philippines	Leonor Cabral-Lim	2003		
Poland	Joanna Jedrzejczak	2001		
Portugal	Isabel Pires*; Joana Parra*; Ines Cunha*; Elia Baeta*; Carla Bentes*; Catarina Cruto*; Inês Menezes Cordeiro*	2001		
Serbia & Montenegro	Maja Milovanovic	2002		
Slovakia	Vladimír Safcák	2002		
Slovenia	Boštjan Čebular & Gal Granda	2002		
Spain	Meritxell Martinez Ferri	2001		
Sweden	Torbjörn Tomson	2000		
Switzerland	Elisabeth Sellitto, Dominique Flügel*	2001		
Taiwan	Hsiang-Yu Yu	2004		
Turkey	Demet Ilhan Algın	2000		
United Kingdom	John Craig & Craig Heath	2001		

NB: Some of the countries listed in this table are currently inactive, not contributing pregnancies the last few years.



By the cut-off date for this report (May 25th, 2023), 29,527 pregnancies had been entered into the central database. Of these, 12,254 pregnancies are excluded from the present interim report for reasons explained here below:

- 1. Pregnancies that failed to meet inclusion criteria (n=217).
- 2. Lost to follow-up, including those failing to submit sub-forms within preset deadlines (n=4,127).
- 3. Pending pregnancies, awaiting updates or corrections of different sub-forms (n=751).
- 4. Ongoing pregnancies, updated and corrected (n=638).
- 5. Retrospective, but completed and corrected (n=4,735). Among these, there are true retrospective pregnancies (n=4,368) and a further three hundred and sixty-seven pregnancies (n=367) that otherwise met our criteria for prospective pregnancies since they were recruited within 16th week, but for which patients had an ultrasound examination performed before enrolment.
- 6. Retrospective, i.e. initially classified as prospective pregnancies but re-classified as retrospective cases because one or more CRF subforms were submitted after the set deadlines (n=407).
- 7. Unclassifiable i.e. cases for which it was impossible to determine if there was a malformation or not (n=93). This includes 1 stillbirth with unknown fetal status, induced abortions with insufficient information on fetus (n=6), anomalies in livebirths where the information was insufficient to determine if qualifying for malformation diagnosis (n=81), 1 incomplete spontaneous abortion with unclear results of biopsy, and 4 perinatal deaths in premature births (<35 gestational weeks) with anomalies difficult to classify as congenital or due to prematurity.
- 8. Not yet classified, i.e. pregnancies which classification is pending as well as pregnancies which became completed after the last time we sent the database to the Outcome Assessment Committee (OAC), regardless if they contained some malformations or not (n=78).
- 9. Treatment changes between different AEDs or mono- to polytherapy or vice versa during the first trimester (n=1,208).

Thus, in total 17,273 prospective pregnancies (enrolled at the latest during the 16th gestational week and before outcome is known) are included in this report.

The classification of the epilepsy among the prospective pregnancies is given in table 2. Epilepsy was the indication for treatment in all but 125 (0.7%) of the pregnant women.

Table 2. Classification of the epilepsy in 17,273 prospective pregnancies.

Epilepsy	N	%
Localisation-related*	9,010	52.2
Generalized	7,207	41.7
Undetermined	583	3.4
Missing information	348	2.0
No epilepsy	125	0.7
Total	17,273	100

^{*}Focal, according to more current terminology.



The women were of Caucasian ethnicity in 86% and of Asian in 10%.

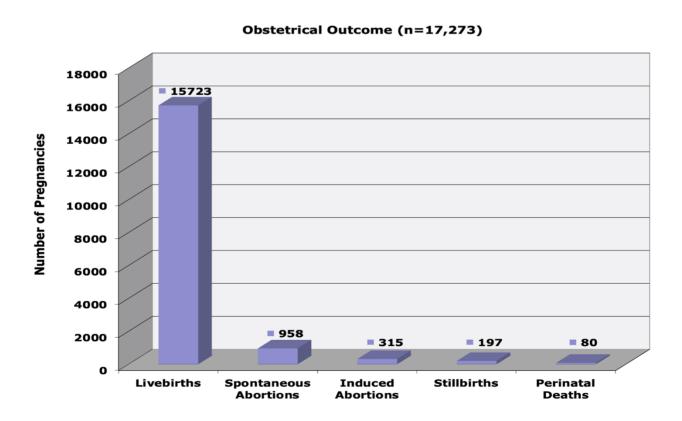
Gravida for each pregnancy is presented in Table 3.

Table 3. Number of the pregnancy in 17,273 prospective cases.

Gravida	N	%
1st pregnancy	7,853	45.5
2nd pregnancy	5,422	31.4
3rd pregnancy	2,390	13.8
4th pregnancy	989	5.7
5th pregnancy	379	2.2
> 5th pregnancy	237	1.4
Not ascertained	3	0.0
Total	17,273	100

The outcome of the prospective completed pregnancies is presented in Figure 2. Out of the 315 induced abortions, 55 were for chromosomal abnormalities and/or syndromes and 82 were for other fetal indication detected by prenatal screening (out of these 82 cases, 69 were confirmed as major malformations and the remaining 13 cases were definitively classified as other abnormalities such as hydrops fetalis, molar pregnancies, blighted ovum, fetal placental transfusion syndromes, fetal growth retardation, fetus papyraceus, fetal death for unspecified causes, balanced translocation and insertion in normal individual).

Figure 2. Obstetrical outcome of prospective pregnancies.

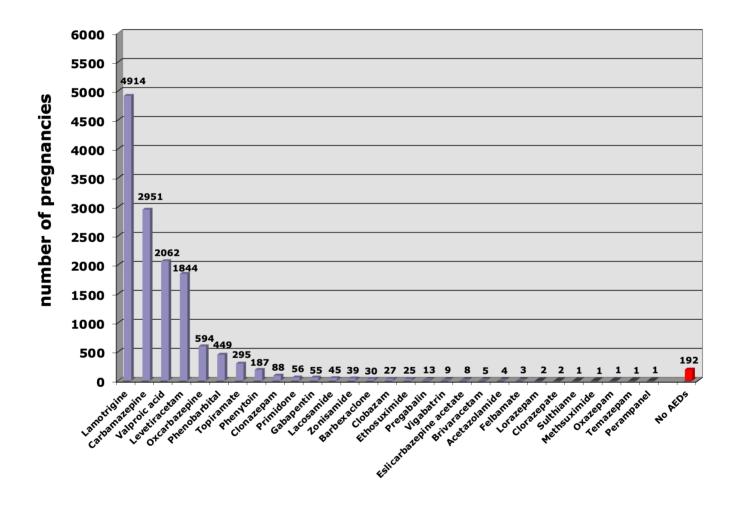




Of the pregnancies, **13,712 (79.4%) involved women on a single AED**, 2,878 (16.7%) were on two AEDs whereas 491 (2.8%) took three AEDs or more. One hundred and ninety-two women (1.1%) were not on AED treatment during the 1st trimester. The exposure to the different AEDs in monotherapy among the prospective pregnancies is presented in Figure 3.

<u>Figure 3</u>. Number of prospective pregnancies with exposure to different AEDs in monotherapy during the first trimester of pregnancy.

Monotherapies (n=13,712)





There were 361 different AED combinations. The most frequently used combinations were lamotrigine and levetiracetam (n=496), lamotrigine and valproic acid (n=301), carbamazepine and levetiracetam (n=188), carbamazepine and clobazam (n=132), carbamazepine and lamotrigine (n=129), lamotrigine and topiramate (n=107), carbamazepine and valproic acid (n=85), carbamazepine and phenobarbital (n=84), clobazam and lamotrigine (n=72), levetiracetam and oxcarbazepine (n=71), levetiracetam and valproic acid (n=67), carbamazepine and topiramate (n=61), and clonazepam and lamotrigine (n=59) (Table 4).

Table 4. The most common AED combinations.

The most common polytherapies during	N
the first trimester of pregnancy	
lamotrigine + levetiracetam	496
lamotrigine + valproic acid	301
carbamazepine + levetiracetam	188
carbamazepine + clobazam	132
carbamazepine + lamotrigine	129
lamotrigine + topiramate	107
carbamazepine + valproic acid	85
carbamazepine + phenobarbital	84
clobazam + lamotrigine	72
levetiracetam + oxcarbazepine	71
levetiracetam + valproic acid	67
carbamazepine + topiramate	61
clonazepam + lamotrigine	59
lacosamide + levetiracetam	48
lamotrigine + oxcarbazepine	47
phenobarbital + valproic acid	41
topiramate + valproic acid	41
clonazepam + valproic acid	40
levetiracetam + topiramate	38
carbamazepine + clonazepam	35
clobazam + oxcarbazepine	35
phenobarbital + phenytoin	33
lamotrigine + phenobarbital	27



The number of pregnancies with exposure to different second generation AEDs taken in combination with other AEDs are listed in Table 5.

<u>Table 5</u>. Number of pregnancies with different second generation AEDs in combination therapy.

Lamotrigine	1,595
Levetiracetam	1,244
Topiramate	422
Oxcarbazepine	298
Lacosamide	125
Zonisamide	115
Gabapentin	66
Vigabatrin	37
Pregabalin	34
Eslicarbazepine acetate	27
Perampanel	27
Brivaracetam	15
Tiagabine	11
Rufinamide	4
Retigabine	1



TERATOGENIC OUTCOME

There were 753 major congenital malformations (MCM), 29 syndromic and/or genetic cases and 97 chromosomal abnormalities (CHR) in the prospective cohort of 16,315 pregnancies as shown in Table 6 (958 spontaneous abortions are excluded).

Table 6. Pathological outcomes.

Outcome	Outcome Classification	N
MCM	Multiple major	62
	Isolated major	691
MCM		753
SYNDROMES or GENETIC conditions		29
CHR		97
Total		879

The 29 syndromic and/or genetic cases are Marfan's syndrome (3), Noonan syndrome (3), inherited tuberous sclerosis (6), Goldenhar syndrome (1), incontinentia pigmenti n.o.s (1), incontinentia pigmenti (Bloch-Sulzberger syndrome) (1), inherited congenital glaucoma (1), inherited congenital cataract (1), inherited craniosynostosis (1), Di George's syndrome (1), bilateral hearing loss (1), X-linked lissencephaly (1), skeletal dysplasia/dwarfism (1), X-linked ichthyosis (1), Freeman Sheldon syndrome (1), Zellweger syndrome (1), achondroplasia (1), blepharophimosis-ptosis-epicanthus syndrome (BPES) (1) and Dravet syndrome (2).

In this report we will confine our analysis to the 753 MCM including 69 induced abortions, seven stillbirths and 18 neonatal deaths. Of the 659 live births, 94 cases of malformations were ascertained prenatally, 384 were first reported at birth, and a further 181 cases not detected at birth but within one year after birth.

Among the 753 cases with MCM, 174 were detected by ultrasound examination. Out of these 174 cases, there were 69 induced abortions, five stillbirths, six perinatal deaths and 94 live births.

The 753 cases represent a **malformation prevalence of 4.6%** of all prospective pregnancies for which follow-up has been completed (753/16,315).



The type of malformations is described in Table 7.

Table 7a - MCMs

## ATHOLOGICAL OUTCOMES MCM			
MCM Multiple major		DESCRIPTION	N
MCM Spins Billids Nervous system MCM Anencephalus and similar MCM Intercephaly MCM Artis spall defect Cardiovascular system MCM Artis spall defect Gardiovascular system MCM Intercephaly MCM Intercephalus MCM Intercephalus	PATHOLOGICAL		
MCM	OUTCOMES		
MCM	MCM	Multiple major	62
MCM Spins affolds 4 MCM Anencephalus and similar 4 MCM Hydrocephaly 4 MCM Microcephaly 3 MCM Nercoes system (other malformations) 1 MCM Arrial septal defect 8 MCM Ventricular septal defect 9 MCM Artivovantricular septal defect 6 MCM Artivovantricular septal defect 6 MCM Congenital hards disease 6 MCM Congenital hards disease 6 MCM Intensity year set westel (complete) MCM Intensity year set set westel (complete) MCM Intensity year set			
MECM Hydrocephaly MCM Microcephaly MCM Nervous system (other malformations) 1 MCM Avial septial defect 3 MCM Avial septial defect 3 MCM Arrivemental defect 3 MCM Congenital heart disease 6 MCM Totalogy of Fallot 1 MCM Transposition of great vessels (complete) 3 MCM Pulmonary valve stenosis 1 1 MCM Pytophastics list he heart 1 1 MCM Phytophastics list he heart 3 1 MCM Phytophastics list he heart 3 1 MCM Interview system 3 6 MCM Interview system 3 6 MCM Interview system 3 6 MCM Displayagnatic hemia 1 6 MCM Displayagnatic hemia 1 1 MCM Displayagnatic hemia 1 1	МСМ		42
MCM Microcosphaly 1 MCM Nervous system (other malformations) 1 MCM Atrial septal defect 3 MCM Atrial septal defect 6 MCM Consental reactions 6 MCM Impropriate the search (complete) 1 MCM Impropriate the search (complete) 2 MCM Dispense search (complete) 2 MCM Dispense search (complete) 2 MCM Dispense search (complete) 2 <td>МСМ</td> <td>Anencephalus and similar</td> <td>5</td>	МСМ	Anencephalus and similar	5
MCM	MCM	Hydrocephaly	7
MCM Atrial septal defect	мсм	Microcephaly	2
MCM Atrial septal defect 3 MCM Ventricular septal defect 6 MCM Atrioueristicular septal defect 6 MCM Attrial septal defect 6 MCM Congenital heart disease 6 MCM Congenital heart disease 6 MCM Tetralogy of sallot 7 MCM Pulmonary valve stenois 7 MCM Unitary system (other malformations) 5 MCM Disphragmatic hemia 7 MCM Disphragmatic hemia 8 MCM Disphragmatic hemia 9 MCM Disphragmatic hemia 9 MCM Disphragmatic hemia 10 MCM Upper imb reduction 10 MCM Upper imb reduction 10 MCM Upper imb reduction 10 MCM Disphragmatic hemia 10 MCM Disphr	MCM	Nervous system (other malformations)	17
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MCM Congenital heart disease 6 MCM Tetratogy of Failot 6 MCM Transposition of great vessels (complete) 1 MCM Pulmonary valve stenosis 1 MCM Hypoplastic left heart 1 MCM Urinary system (other malformations) 5 MCM Lenal Dysplasia a) 6 MCM Internal Dysplasia 6 6 MCM Dispestive system 6 6 MCM Dispestive system 7 7 MCM Dispestive system (other malformations) 1 1 MCM Objective system (other malformations) 1 1		·	69
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MCM Duodenal atresia or stenosis MCM Gastrocchisis MCM Omphalocele MCM Atresia of oesophagus without fistula all 3 MCM Upper limb reduction MCM Lower limb reduction MCM Lower limb reduction MCM Syndactyly 1 MCM Syndactyly 2 MCM Club foot - tallpase equinovarus 2 MCM Club foot - tallpase equinovarus 2 MCM Limbs (other malformations) 3 MCM Musculo-skeletal (other malformations) 1 MCM Musculo-skeletal (other malformations) 1 MCM Hip dislocation and/or dysplasia 8 MCM Hypospadias 8 MCM Hypospadias 8 MCM Developmental ovarian cyst MCM Genital (other malformations) 8 MCM Genital (other malformations) 1 MCM Congenital cataract 1 MCM <td>MCM</td> <td>Ano-rectal atresia and stenosis</td> <td>2</td>	MCM	Ano-rectal atresia and stenosis	2
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MCM Upper limb reduction MCM Lower limb reduction MCM Syndactyly MCM Polydactyly MCM Polydactyly MCM Club foot- talipes equinovarus MCM Limbs (other malformations) MCM Musculoskeletal MCM Musculoskeletal MCM Musculoskeletal MCM Hip dislocation and/or dysplasia MCM Method Musculoskeletal (other malformations) MCM Hypospadias MCM Method M			
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MCM Lower limb reduction MCM Syndactyly MCM Polydactyly 2 MCM Club foot - talipes equinovarus 2 MCM Limbs (other malformations) 3 MCM Musculo-skeletal (other malformations) 1 MCM Musculo-skeletal (other malformations) 1 MCM Hip dislocation and/or dysplasia 7 MCM Hip obspadias 8 MCM Developmental ovarian cyst 8 MCM Genital (other malformations) 8 Eye, ear, face and neck MCM Congenital cataract 9 MCM Eye (other malformations) 1 MCM Eye, (other malformations) 1 MCM Atresia of nasopharynx all 1 MCM	NACNA	- i	
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MCM Limbs (other malformations) Musculo-skeletal (other malformations) 1.			23
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MCM Developmental ovarian cyst		all	86
MCM Genital (other malformations) all 8i Eye, ear, face and neck MCM Congenital cataract MCM Eye (other malformations) MCM Eye, ear neck MCM Eye, ear neck MCM Eye, ear neck MCM Choanal atresia MCM Choanal atresia MCM Atresia of nasopharynx all 1: Oro facial clefts MCM Cleft lip with or without palate MCM Cleft palate Other specified malformations (including sacral teratoma, cystic hygroma, haemangiomas, accessory skin tags, aberrant subclavian artery, congenital malformation of renal artery, congenital malformation of adrenal gland,congenital malformations of integument, congenital malformations of the lung, congenital bronchomalacia, congenital malformations of thyroid gland). MCM all MCMs Syndromes all Syndromes		Genital system	
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MCM all MCMs 753 CHR all CHR 97 Syndromes all Syndromes 29			
CHR all CHR 97 Syndromes all Syndromes 29			31
Syndromes all Syndromes 29	MCM	all MCMs	753
	CHR	all CHR	97
	Syndromes	all Syndromes	29
	Total	all cases with pathological outcomes	879



Table 7b – CHR & Syndromes

PATHOLOGICAL	ATHOLOGICAL DESCRIPTION		
OUTCOMES			
MCM	all MCMs	753	
	Chromosomal		
CHR	Chromosomal	24	
CHR	Down's syndrome	48	
CHR	Edward syndrome/trisomy 18	10	
CHR	Klinefelter's syndrome	2	
CHR	Patau syndrome/trisomy 13	6	
CHR	Turner's syndrome	5	
CHR	Wolff-Hirschorn syndrome	2	
CHR	all CHR	97	
	Syndromes or genetic conditions		
Syndrome	Marfan's syndrome	3	
Syndrome	incontinentia pigmenti, n.o.s	1	
Syndrome	incontinentia pigmenti (Bloch-Sulzberger syndrome)	1	
Syndrome	Noonan's syndrome	3	
Syndrome	Goldenhar syndrome (oculo-auriculo-vertebral syndrome)	1	
Syndrome	Di George's syndrome	1	
Syndrome	tuberous sclerosis	6	
Syndrome	craniosynostosis, inherited	1	
Syndrome	congenital cataract, inherited	1	
Syndrome	congenital glaucoma, inherited	1	
Syndrome	X-linked ichthyosis	1	
Syndrome	X-linked lissencephaly	1	
Syndrome	hearing loss, bilateral, inherited	1	
Syndrome	skeletal dysplasia (achondroplastic dwarfism)	1	
Syndrome	Freeman Sheldon Syndrome (distal arthrogryposis type 2A)	1	
Syndrome	Zellweger syndrome	1	
Syndrome	achondroplasia	1	
Syndrome	blepharophimosis-ptosis-epicanthus syndrome (BPES syndrome)	1	
Syndrome	Dravet syndrome	2	
Syndromes	all Syndromes	29	
Total	all cases with pathological outcomes	879	



In 553 out of 12,989 pregnancies with AED monotherapy, one or more MCMs were observed (4.3%) as opposed to 194 out of 3,140 pregnancies with AED polytherapy (6.2%), as shown in Table 8.

<u>Table 8</u>. Pathological outcomes by AED treatment categories.

(In this table, 958 spontaneous abortions have been excluded from the denominator).

	No AED	%	Monotherapy	%	Polytherapy	%	Total
MCM	6	3.2	553	4.3	194	6.2	753 (4.6%)
CHR	2	1.1	78	0.6	17	0.5	97 (0.6%)
Syndromes	0	0.0	23	0.1	6	0.2	29 (0.2%)
No malformation	178	95.7	12,335	95.0	2,923	93.1	15,436 (94.6%)
Total	186	100	12,989	100	3,140	100	16,315 (100%)

PUBLICATIONS

Changes in AED prescribing patterns and in rates of MCM over time in the EURAP cohort were published in *Neurology*. 2019 Aug 27;93(9):e831-e840.

Outcome regarding the eight most common monotherapies has been published in *Lancet Neurology, April* 18, 2018.

The dose-dependent risk of MCM with exposure to valproate in mono- and polytherapy has also been analysed and reported (*Neurology*, *Sept 8*, *2015*) and so has the risk of intrauterine death in association with different treatments (*Neurology Aug 18*, *2015*).

A manuscript on seizure control in pregnancies with withdrawal of or switch from valproate during 1st trimester as compared with maintained valproate treatment has been published in Epilepsia (*Epilepsia 2016;* 57: e173-7).

Outcome in relation to exposure to individual drugs or specific drug combinations is not included in the present report.

ORGANISATION, FUNDING AND SUPPORT

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APPENDIX

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