

**PODRŽITE OSOBE S EPILEPSIJOM -
NOSITE LJUBIČASTO!**

**26. OŽUJKA
LJUBIČASTI DAN**



"Imaš epilepsiju? Znaj da nisi sam!"

- Cassidy Megan, osnivačica Ljubičastog dana

**Obilježavanje
Ljubičastog dana
u KBC-u Zagreb
25.03.2021.**

Purple Day

- Cilj: edukacija obitelji, nastavnika, a posebno djece školske dobi kako bi zauzeli što pozitivniji stav prema epilepsiji u budućem životu
- Destigmatizacija
 - značajni pomaci od 2008. god.
- pomoći oboljelima da ih okolina prihvati i razumije njihovu bolest



Što je to **EPILEPSIJA?**

poremećaj funkcije mozga koji se
očituje epileptičnim napadajima

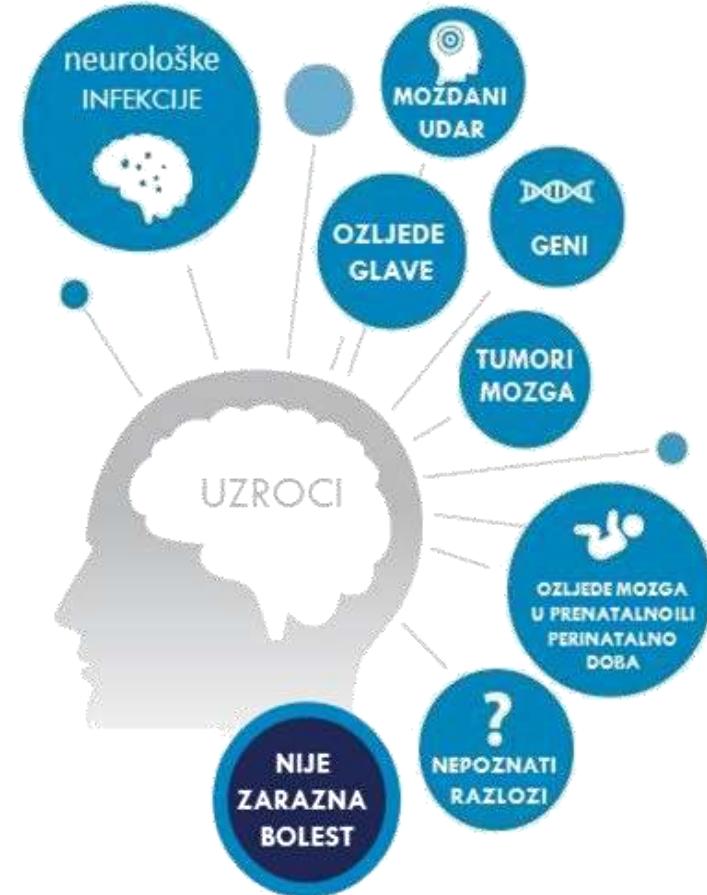
50 000 000 ljudi

Širom svijeta ima epilepsiju

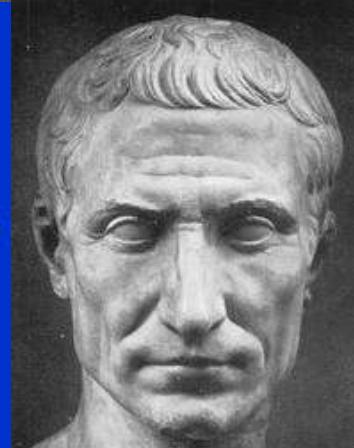
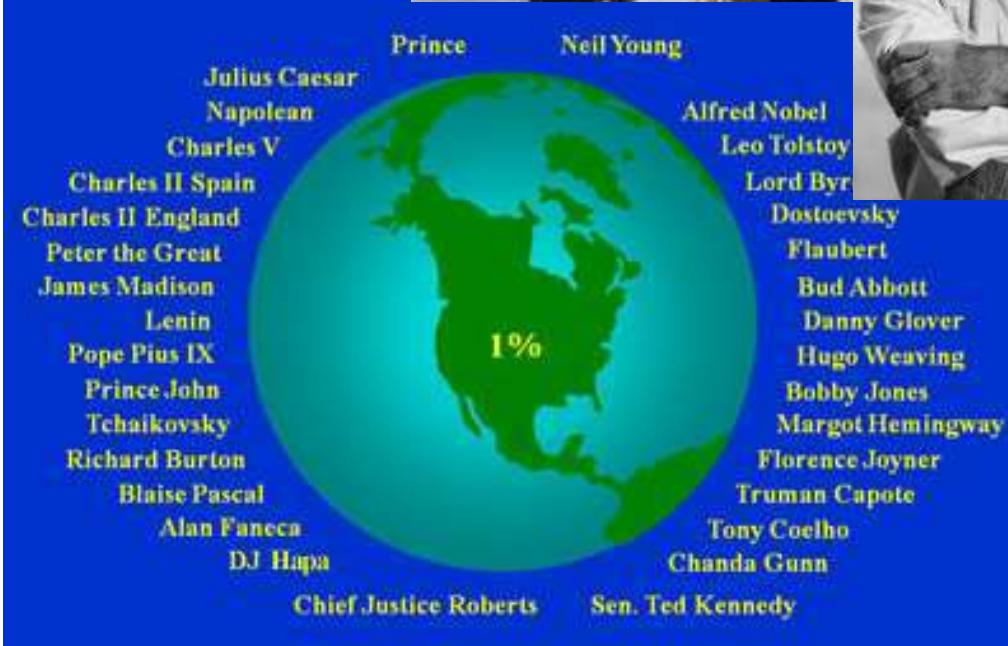
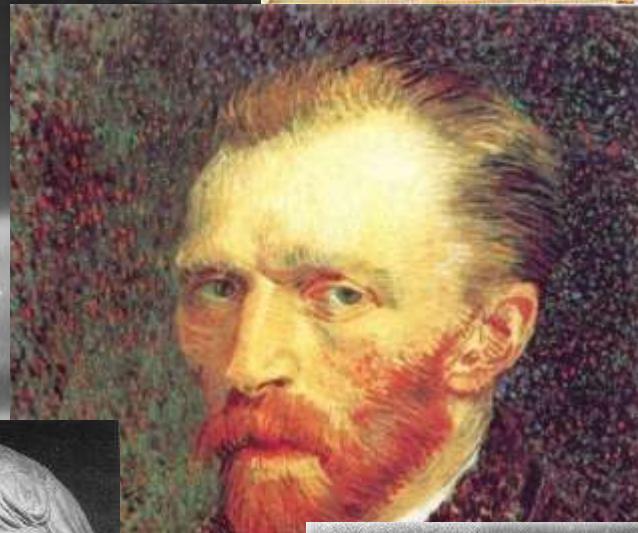
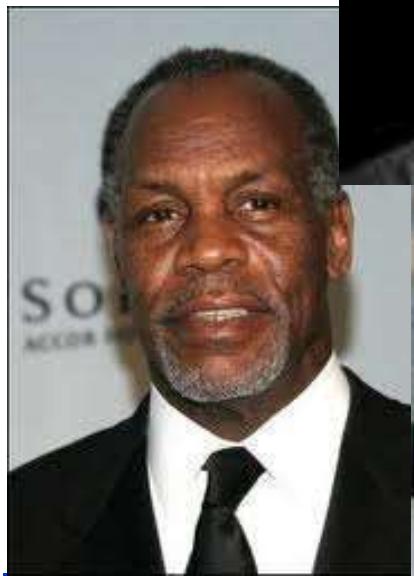


40 000 oboljelih u Hrvatskoj

Izvor: Global Burden of Disease, SZO



Sistemske poremećaje: hipoglikemija, hiponatrijemija, uremija, hepatička encefalopatija, intoksikacije, hipertermija, eklampsija

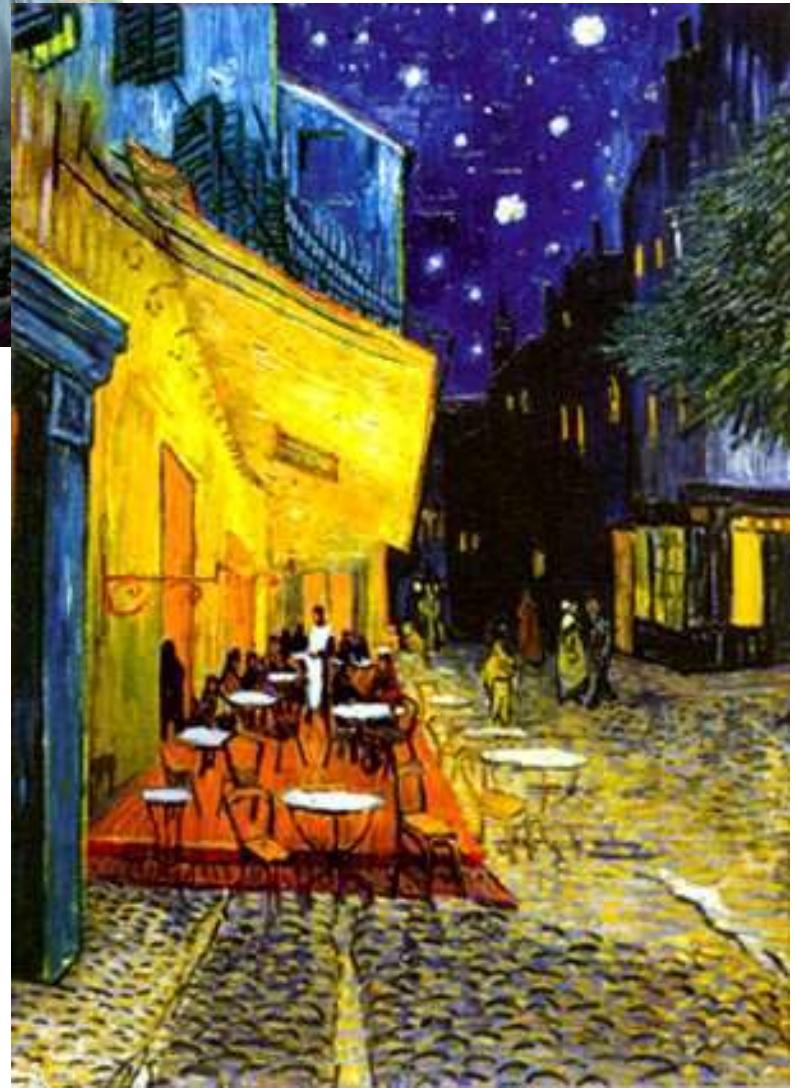




“Alice's Adventures in Wonderland”

Lewis Carroll (1832 -1898)

Vincent Van Gogh: “Cafe Terrace by Night”, 1888. god



Dijagnoza epilepsije

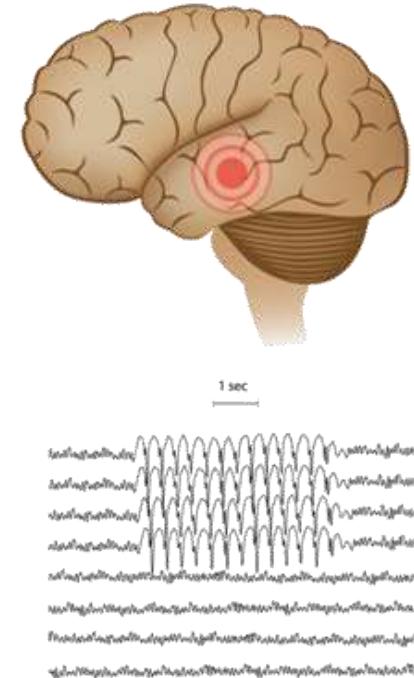
1. Najmanje dva neprovocirana epileptična napadaja koja su se dogodila u vremenskom razmaku duljem od 24 sata.
2. Jedan neprovocirani epileptični napadaj i vjerojatnost dalnjih napadaja najmanje 60% u narednih 10 godina (bolesnici s preboljelim moždanim udarom, infekcijom središnjeg živčanog sustava, kod promjena u EEG-u, kod određenih tipova traume mozga, tumora mozga, itd.)
3. Dijagnoza epileptičnog sindroma

A practical clinical definition of epilepsy

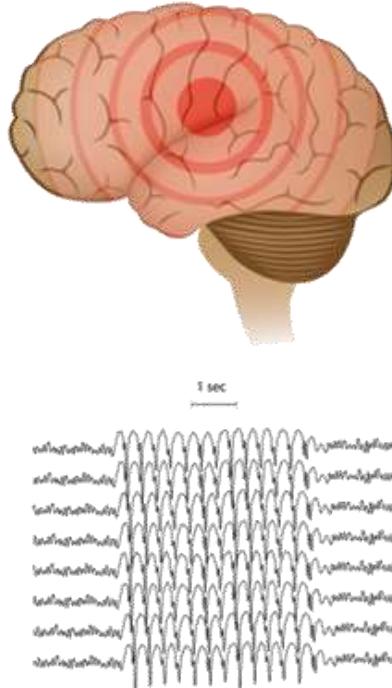
*Robert S. Fisher, †Carlos Acevedo, ‡Alexis Arzimanoglou, §Alicia Bogacz, ¶J. Helen Cross,
#Christian E. Elger, **Jerome Engel Jr, ††Lars Forsgren, ‡‡Jacqueline A. French, §§Mike
Glynn, ¶¶Dale C. Hesdorffer, ##B.I. Lee, ***Gary W. Mathern, †††Solomon L. Moshé,
‡‡‡Emilio Perucca, §§§Ingrid E. Scheffer, ¶¶¶Torbjörn Tomson, #####Masako Watanabe, and
****Samuel Wiebe

Klasifikacija epilepsija

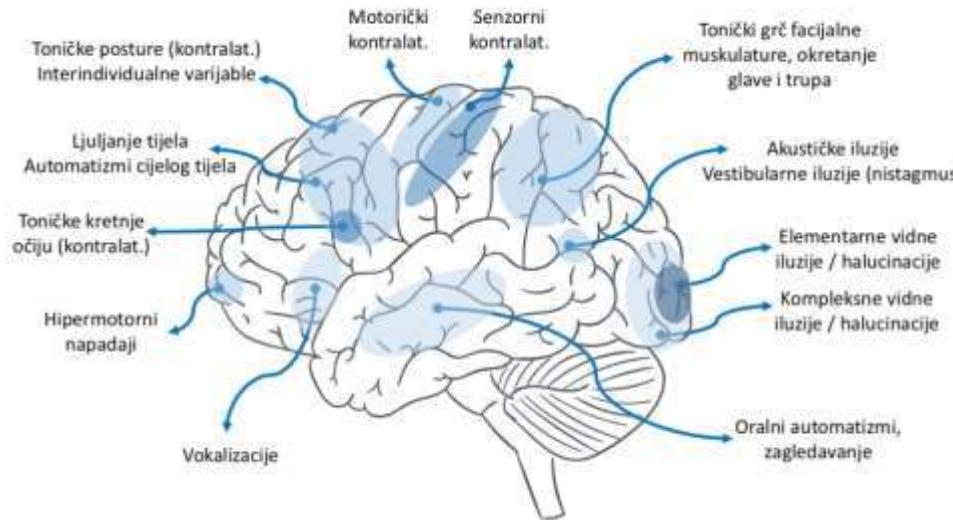
1. Žarišne epilepsije



2. Generalizirane epilepsije



Klinička semiologija napadaja



3. Epilepsije nepoznatog početak

Specijalni epileptični sindromi



PATOFIJOLOGIJA EPILEPSIJE

paroksizmalni poremećaj funkcije mozga koji je rekurentan i stereotipan



stanice moždane kore - previše podražljive, reagiraju s ekscesivnim, sinkronim i, u početku, ograničenim kortikalnim neuronalnim izbijanjima

klinički - epileptični napadaji



zagledavanje

nevoljne, nesvrishodne kretnje

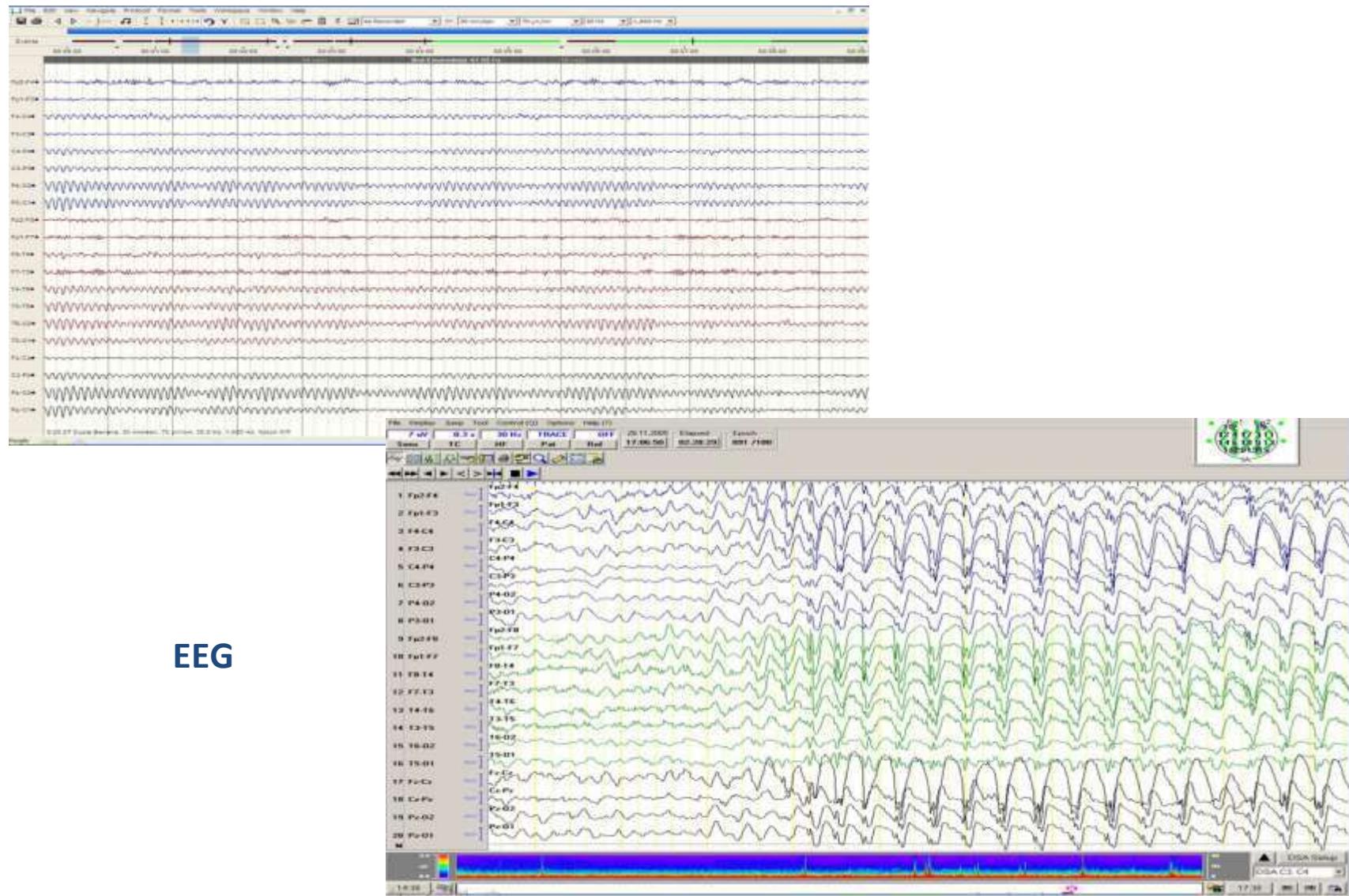
trzajevi

zbunjenost

VRSTE EPILEPTIČNIH NAPADA

Dijagnostika epilepsije





Postoji li RJEŠENJE ?



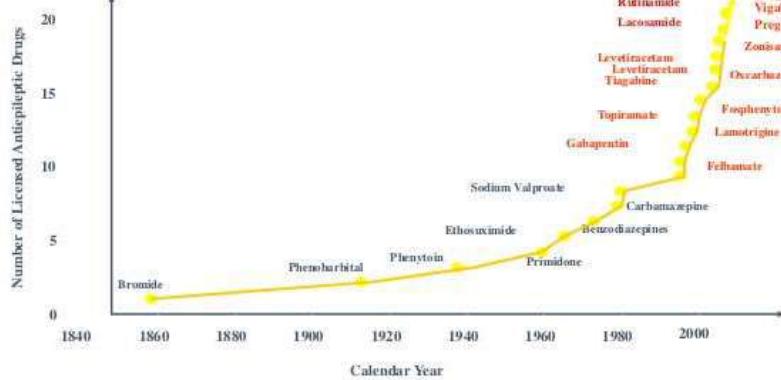
Epilepsija se može liječiti djelotvornim, povoljnim i dostupnim antiepilepticima

70%
NORMALAN ŽIVOT



S redovitom terapijom 70% oboljelih od epilepsije može ostvariti punu kvalitetu života

ANTIEPILEPTIC DRUG DEVELOPMENT

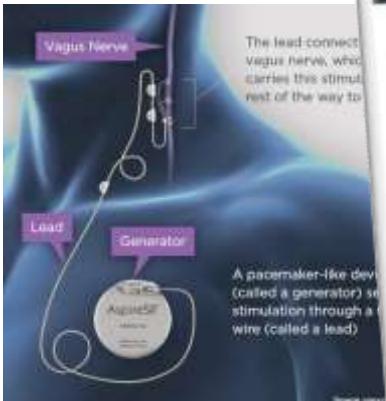
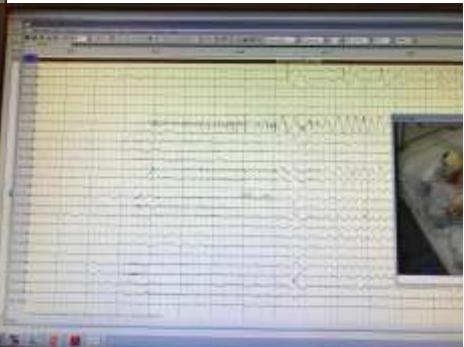
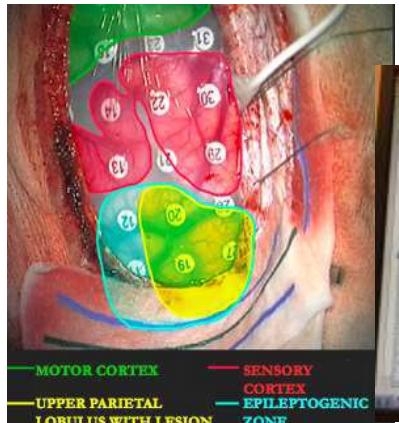


„Klasični“ (stari) antiepileptici:

fenobarbiton (PB)
metilfenobarbiton (MPB)
fenitojn (PHT)
acetazolamid (ACT)
primidon (PRM)
etosuksimid (ESC)
sultiam (SUL)
karbamazepin (CBZ)
valproati (VPA)
klonazepam (CNZ)
klobazam (CLB)
piracetam (PCT)

„Moderni“ (novi) antiepileptici:
vigabatrin (VGB)
lamotrigin (LTG)
gabapentin (GBP)
felbamat (FBM)
topiramat (TPM)
tiagabin (TGB)
okskarbazepin (OXC)
levetiracetam (LEV)
pregabalin (PGB)
zonisamid (ZNS)
rufinamid (RUF)
stiripentol (STP)
lakozamid (LCS)
eslikarbazepin-acetat (ESL)
retigabin (RTG)
perampanel (PER)
brivaracetam (BRV)

Kirurško liječenje epilepsije

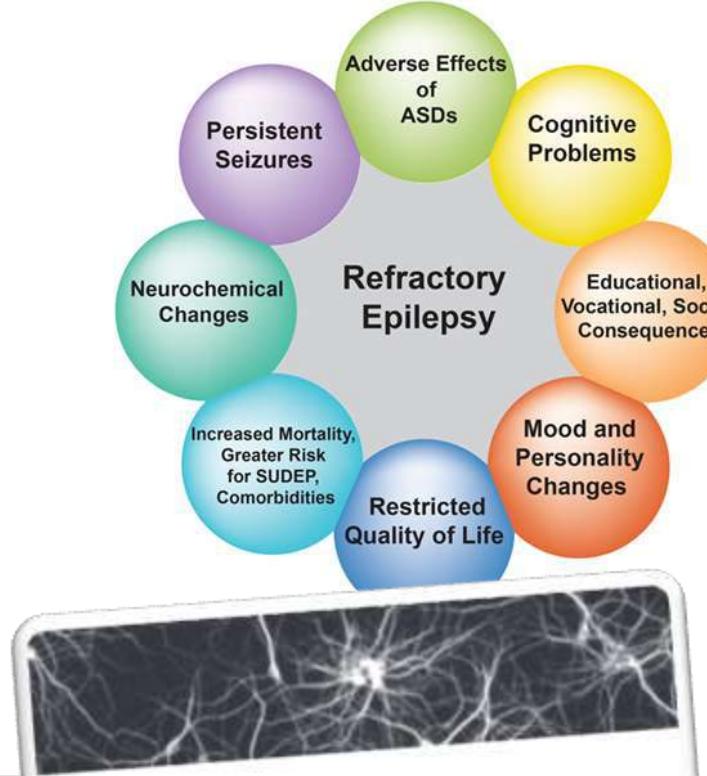


Prvo invazivno monitoriranje i operacija mozga bolesnice s farmakorezistentnom epilepsijom u Hrvatskoj

Dana 23. travnja 2018. godine prvi put u Hrvatskoj u ženskoj regiji bolesnica je u KBC-u Zagreb na Šubiću, uđeno operacije 46 godišnje bolesnici s teškotinom farmakorezistentne oblike epilepsije, koju su imao od 7. godine života i u kojoj, usprkos četiri razne antiepileptičke terapije, nisu postignuti zadovoljavajući lečenijski rezultati. Epilepsija je počela u dobi 20-tak godina, započela se u danu, žestu u mimo, blizinski fizijski radilo se u vremenu, kognitivnoj funkciji i funkciji života dobro je izložena. Iako je u prethodnoj operaciji neurocirurgičkom remozirano mnogo loci, te izradeno o komplikaciji i drugim, nije učinkovito predstavljanje Klinike za neurologiju do sreće. Sve je potencijalno moglo dovesti do posljedice dobiti. Prof. dr. sc. Josipa Poljaković, u svrhu prepreka ovih posljedica, u sklopu kliničke akademije Željka Bratčića i prof. dr. sc. Ivica Wölfle, predmetnoj klinici na epilepsiju hrvatskih bolnika u Beogradu, uspostavio je tada Željkova rekonstrukcijska klinika, dr. sc. Svetozara Poljakovića, dr. Marija Matović, dr. sc. Ivica Škoko, dr. sc. Tatjana Petrić, dr. sc. Željka Poljaković, dr. Štefka Novaković, dr. Vesna Škoko, dr. sc. Božidar Šarić, neurokirurg dr. sc. Željko Pašalić, neurofisiolog dr. sc. Goran Mirković, dr. sc. Andrej Drmotačić, te dr. sc. Goran Mirković, dr. sc. Andrej Drmotačić.



Sl. 1. Tima koji je učestvovao u invazivnom monitoriranju i sljedećoj operaciji bolesnice s farmakorezistentnom oblikom epilepsije u Hrvatskoj i u Šubiću (leži: dr. sc. Goran Mirković, dr. sc. Željko Poljaković, dr. sc. Štefka Škoko, dr. sc. Tatjana Petrić, dr. sc. Ivica Škoko, dr. sc. Željko Poljaković, dr. Štefka Novaković, dr. Vesna Škoko, dr. sc. Božidar Šarić, neurokirurg dr. sc. Željko Pašalić, neurofisiolog dr. sc. Goran Mirković, dr. sc. Andrej Drmotačić).



**Referentni centar Ministarstva
zdravstva RH za epilepsiju
Affiliated to ERN EpiCARE
Klinika za neurologiju Medicinskog
fakulteta u Zagrebu i KBC Zagreb**



Prof. dr. sc. Franjo Hajnšek



Prof. dr. sc. Nikola Gubarev



EEG laboratorij 1958. god.

i

2021. god.

50 godina Centra za epilepsiju KBC-a Zagreb

• U prosljednjima Novitarskog dana u Zagrebu 18. je prosljena svečana objetna 50. obljetnica Centra za epilepsiju KBC-a Zagreb, vodećeg središta ne samo u Hrvatskoj, već i regiji za dijagnostiku i liječenje ove neurološke bolesti. U organizaciji Klinike za neurologiju KBC-a Zagreb i Referentnog centra za epilepsiju Ministarstva zdravstva i socijalne skrbi te pod pokroviteljstvom Ministarstva zdravstva i socijalne skrbi RH, ravnatelja KBC-a Zagreb akademika Željka Belinera, Grada Zagreba, Gradskog ureda za zdravstvo i branitelje, te zagrebačkog Medicinskog fakulteta, skupu su prisustvovali svjetski poznati neurolozi te mnogi vršnici stranci.

Stručnjaci iz Centra su svojim predavanjima predstavili novodobne dijagnostičke mogućnosti i naprednu postoljnici u liječenju epilepsije (medikamentozno, kliničko rezekcionalno i menentreno) i invazivnoj neurokirurgiji (operaciji implantacijom vagusovih stimulatora). Uvodnicu nuc obuzeo je ravnatelj KBC-a Zagreb, akademik Željko Beliner, a izlaganja su potom nastavili doc. dr. Sanja Hajnsek, doc. dr. Željka Petelin Gadžić, dr. Boško Rastovčan i dr. Katarina Blazina.

Centar za epilepsiju osnovao je prof. dr. Franjo Hajnsek 1958. godine. Do stamnositelja hrvatske bio je referentno jedinice na razini bivše Jugoslavije, a od 1999. postaje zauzajom prof. dr. Nikole Gobareva Referentni centar za epilepsiju Ministarstva zdravstva i socijalne skrbi. Doc. dr. Šanja Hajnsek, pročelnica Centra za epilepsiju i predstavnica Klinike za neurologiju KBC-a Zagreb, u svom je obraćanju, među ostalim, istaknula:

„Centar za epilepsiju KBC-a Zagreb u prošlim 50 godinama konstantno je prati aktualne svjetske trendove te pacientima u Hrvatskoj i regiji osigurava vrhunsku razinu liječenja i uspehe rezultate kod liječenja ove neurološke bolesti. Zahvalila bih se Upravi KBC-a Zagreb i ravnatelju akademiku Željku Belinoru na razumijevanju za potrebe Centra i pomoći oko nasveje novih najutjecajnijih trendova te svim ostalim institucijama koji su progongili da ovaj Centar bude ne samo

referentno središte u Hrvatskoj, već da ulazi u svjetski epileptološki krugovor. Posedujem sam ponos na svoj tim, jer smo 23. studenog 2009., kao prvi u Hrvatskoj, na pacijentku koja je kandidat za kirurško liječenje izvršili Wada test, svjetski prvi put postupak za utvrđivanje lokacije funkcije govora i pamćenja u dominantnoj, odnosno ne-dominantnoj mještanoj hemisferi.“

Upravo ovaj test, koji su po prvi put u Hrvatskoj izvršili hipofizički - neurologi doc. Hajnsek, doc. dr. Željka Petelin Gadžić i doc. Petelin Gadžić te neuroendokolog dr. David Ozreček - uz magnesiju rezonansnu magnetnu i semi-invazivnu EEG monitorizaciju, kroz se u sklopu preoperativne obrade bolesnika s epilepsijom kod kojih lijekovima nije postignuta zadovoljavajuća kontrola napada. Na unapređivanje dosadašnjih preoperativnih obrada bolesnika i upravo takvom polikliničkom epilepsiju krećemo se kroz suradnju:

3. Kliničkom za epileptologiju Sveučilišta u Bonu; što predstavlja još jedno puštanje ove godine Centra.

U Europi za sada samo dva medicinska centra (u Velikoj Britaniji i u Francusku) koriste Nicolet Koriski Stimulator kao neselosticirajući uređaj takvog tipa, koji pre stavlja napomenu inovaciju u funkcionalno EEG monitoriranju. Zahvaljujući podršci Uprave KBC-a Zagreb do kraja 2009. godine će uređaj dobiti i Centar za epilepsiju Klinike za neurologiju KBC-a Zagreb. Ova uređaj je za nevidljene mogućnosti obavijajući detaju prikaz moždane kartografije (kortikalni centra za motoriku, govor i smjer) te ih određiti epileptičkih napada (epileptičkih fokusu), kinzijskoj sali, kod EEG monitoriranja za injekt resekcije tumora ili kinzijskih zahvata kod epilepsije.

Zbog dugogodišnje vodeće uloge u liječenju i dijagnostici epilepsije, u Centru još uvijek dolaze, osim iz Hrvatske, i pacijenti iz susjednih zemalja. Uz pročelnicu doc. Hajnsek, u Centru je zapošljeno još šest specijalista neurologija: doc. Petelin Gadžić, doc. Poljanović, dr. Šibila Nanković, mr. sc. dr. Borisla Radic, dr. Vlatko Šulentić i dr. Ivana Kovacević - koji sudjeluju u organiziranju i provođenju trajne edukacije specijalista neurologije u području epileptologije. U tim Centru su također uključeni i neurokurg, neurodiplod, poljopristar, specijalist nuklearne medicine, neuropsiholog i socijalni radnik, koji u



Tim Centra za epilepsiju, KBC Zagreb (s lipom na desnoj):

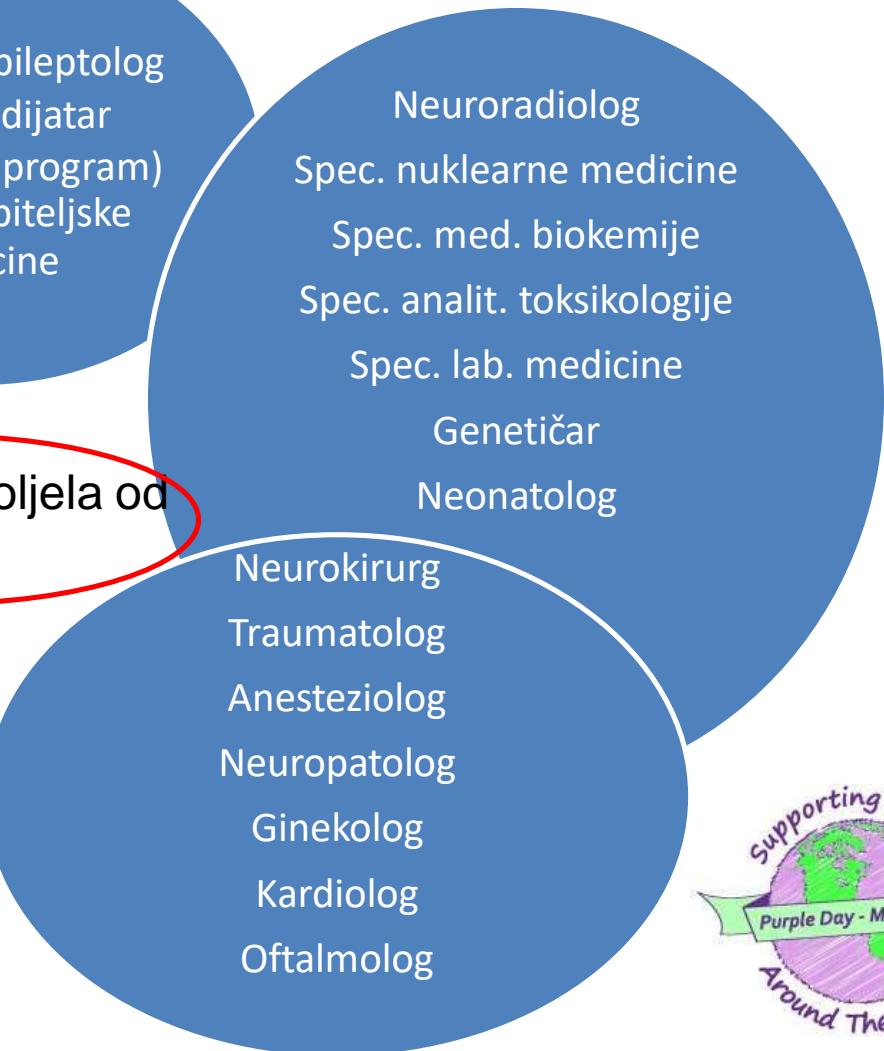
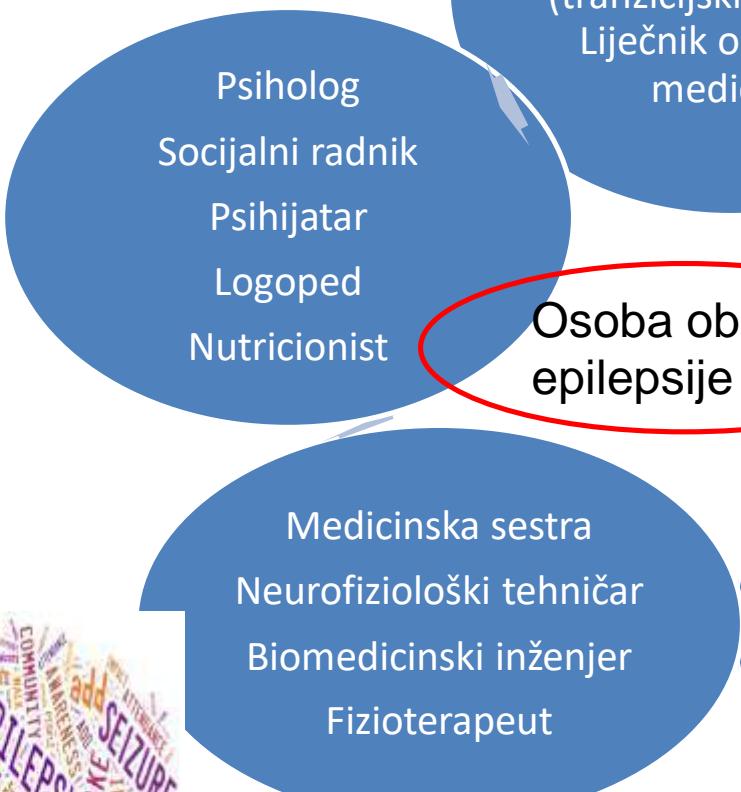
I. Dadić, J. Pečarić, S. Nenčević, J. Bošnjak, R. Novosel, V. Salamović, Š. Hajnsek, M. Jevremović, R. Garićanin, Ž. Petelin Gadžić, I. Kovacević, B. I. Šepet, M. Herderić, B. Radic,



2009. god.



Multidisciplinarni pristup



Suradnja s laboratorijima u Klinici za neurologiju

*Laboratorij za
EMNG*

*Laboratorij za
testiranje
autonomnog
živčanog sustava*

Laboratorij za UZV

*Laboratorij za
kognitivnu i
eksperimentalnu
neurofiziologiju*

*Laboratorij za
evocirane
potencijale*

Dnevna bolnica

University Hospital Centre Zagreb, School of Medicine, University of Zagreb, Croatia

Department of Neurology

Head: Prof. Ervina Bilić, MD, PhD

Epilepsy Centre:

Assoc. Prof. Željka Petelin Gadže, MD, PhD

Prof. Zdravka Poljaković, MD, PhD

Sibila Nanković, MD

Vlatko Šulentić, MD, MSc

Andreja Bujan Kovač, MD

Petra Nimac Kozina, MD

Biljana Đapić Ivančić, MD

Assist. Prof. Magdalena Krbot Skorić, EE, PhD

Valentina Ričković, clinical psychologist

Neurophysiology technicians: Jela Dodig, Maja Jovanović,

Ruža Novosel, Jasna Pekčec, Mirjana Hudoletnjak

Epilepsy Nurses: Nikolina Hederić, Jasminka

Korenika, Anita Ljubić, Nives Šestić

Centre for Translational and Clinical Research:

Prof. Fran Borovečki, MD, PhD

Kristina Gotovac Jerčić, MD, PhD

Antonela Blažeković, MD

Department of Neuroradiology:

Assoc. Prof. Marko Radoš, MD, PhD

Assoc. Prof. Milan Radoš, MD, PhD

Assist. Prof. David Ozretić, MD, PhD

Assist. Prof. Goran Pavliša, MD, PhD

Ivan Jovanović, MD

Department of Nuclear Medicine:

Ratimir Petrović, MD

Anja Tea Golubić, MD

Department of Neurosurgery:

Assist. Prof. Goran Mrak, MD, PhD

Andrej Desnica, MD

Jakob Nemir, MD, PhD

Niko Njirić, MD

Lana Maljković, neuropsychologist

Department of Neuropathology:

Prof. Kamelija Žarković, MD, PhD

Antonija Jakovčević, MD, PhD

Department of Psychiatry and Psychological Medicine:

Assist. Prof. Saša Jevtović, M.D., Ph.D.

Maja Živković, M.D., Ph.D.

Department of Pediatrics:

Prof. Nina Barišić, MD, PhD

Branka Bunoza, MD

Ivan Lehman, MD

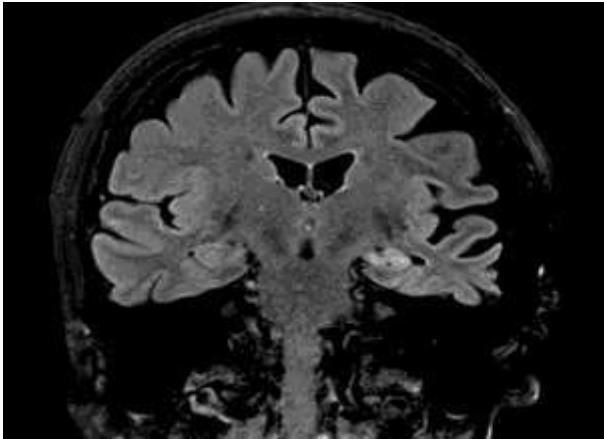
Nataša Nenadić Baranašić, MD, PhD

Department of Laboratory Diagnostics:

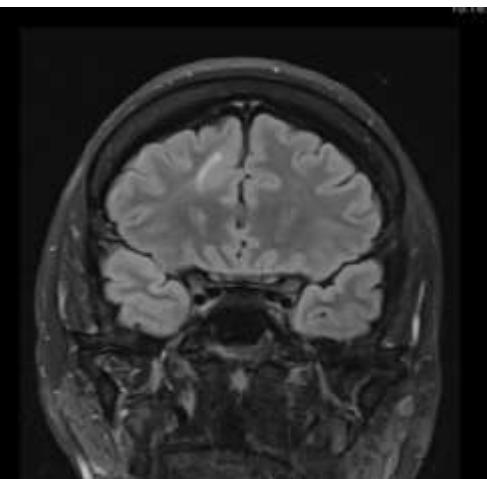
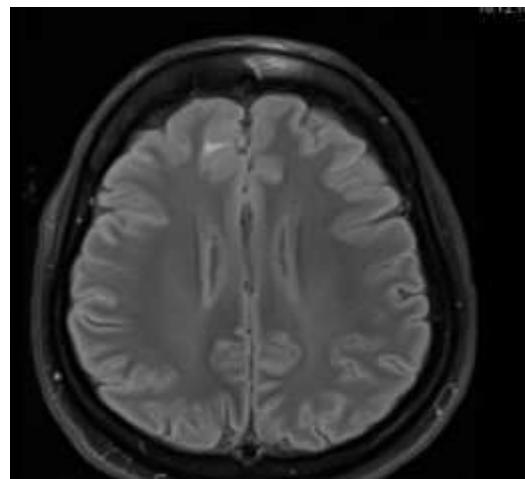
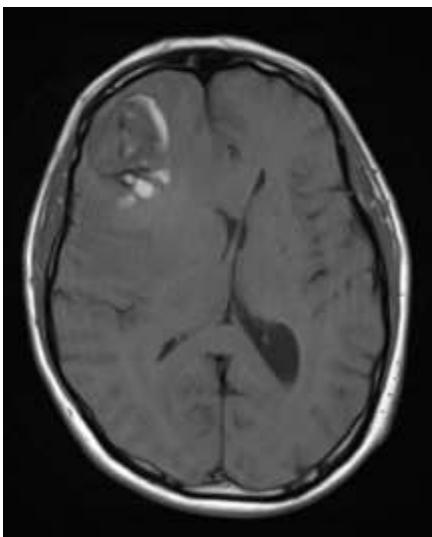
Prof. Nada Božina, MD, PhD

Assist. Prof. Mila Lovrić, PhD

Ana Kozmar, PhD



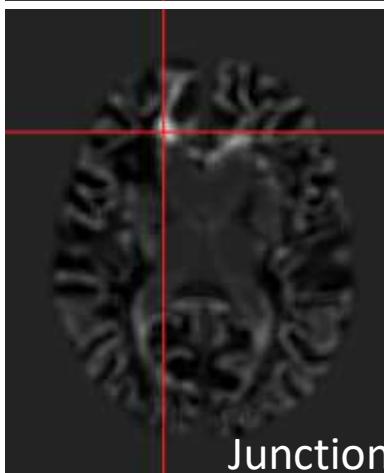
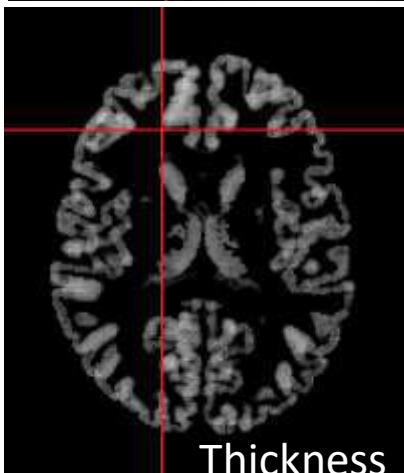
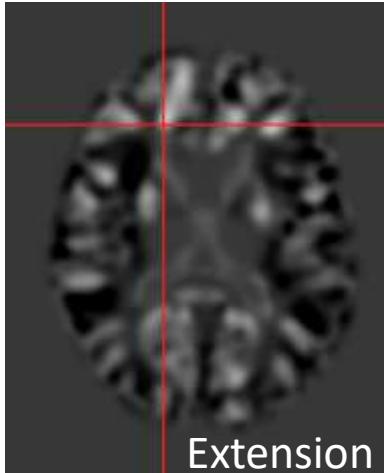
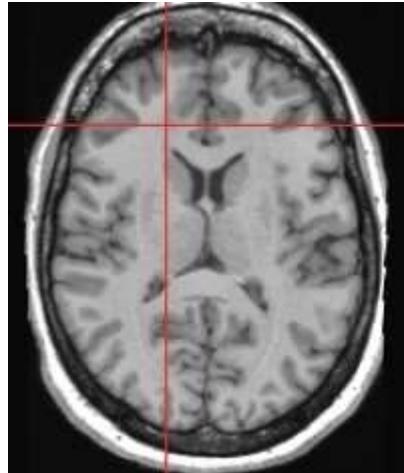
MR mozga





Brain MRI postprocessing software

Detection of FCD and other types of MCD



Brain MRI post-processing with MAP07 in the preoperative evaluation of patients with pharmacoresistant epilepsy – Croatian single centre experience

Andreja Bujan Kovac^{a,b,*}, Zeljka Peselin Gadze^a, Milan Rados^b, Magdalena Krbot Skoric^{a,c}, Goran Mrak^d, Jakob Nemir^d, Milan Mihovcic^e, Sanja Hajnsek^f

^a Department of Neurology, University Hospital Center Zagreb, School of Medicine, University of Zagreb, Referral Centre of the Ministry of Health of the Republic of Croatia for Epilepsy, Affiliated Partner of the HRM Research, Zagreb, Croatia

^b Croatian Institute for Brain Research, School of Medicine, University of Zagreb, Zagreb, Croatia

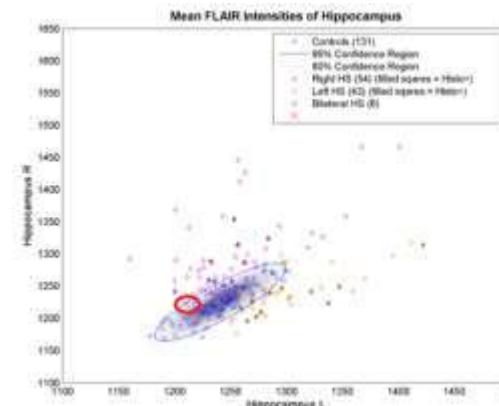
^c University of Zagreb, Faculty of Electrical Engineering and Computing, Zagreb, Croatia

^d Department of Neurosurgery, University Hospital Center Zagreb, School of Medicine, University of Zagreb, Affiliated Institute of HRM/DAK, Zagreb, Croatia

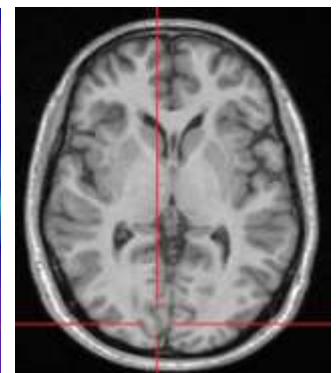
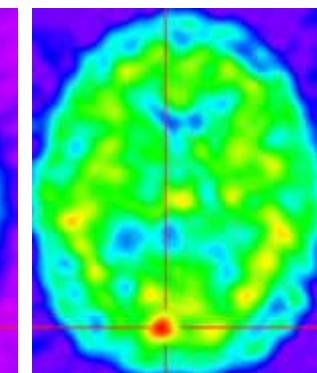
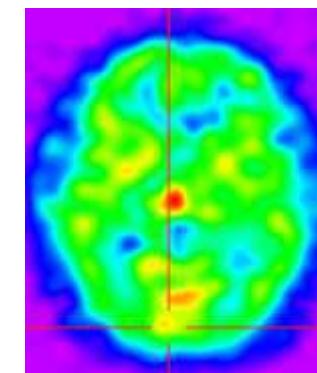
^e Andrija Štampar School of Public Health, Department for Environmental Health, Occupational and Sports Medicine, University of Zagreb, School of Medicine, Zagreb, Croatia

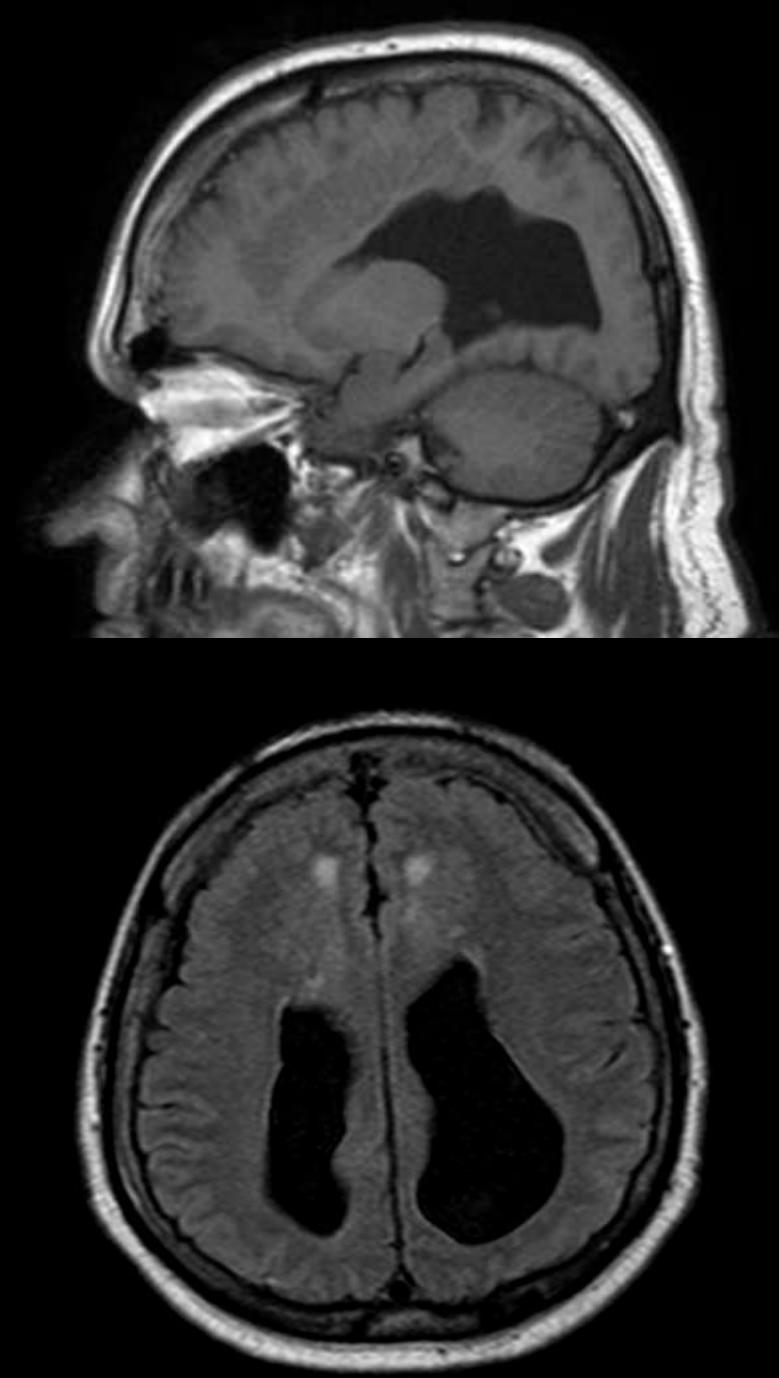
^f School of Medicine, University of Zagreb, Zagreb, Croatia

Detection of MTS

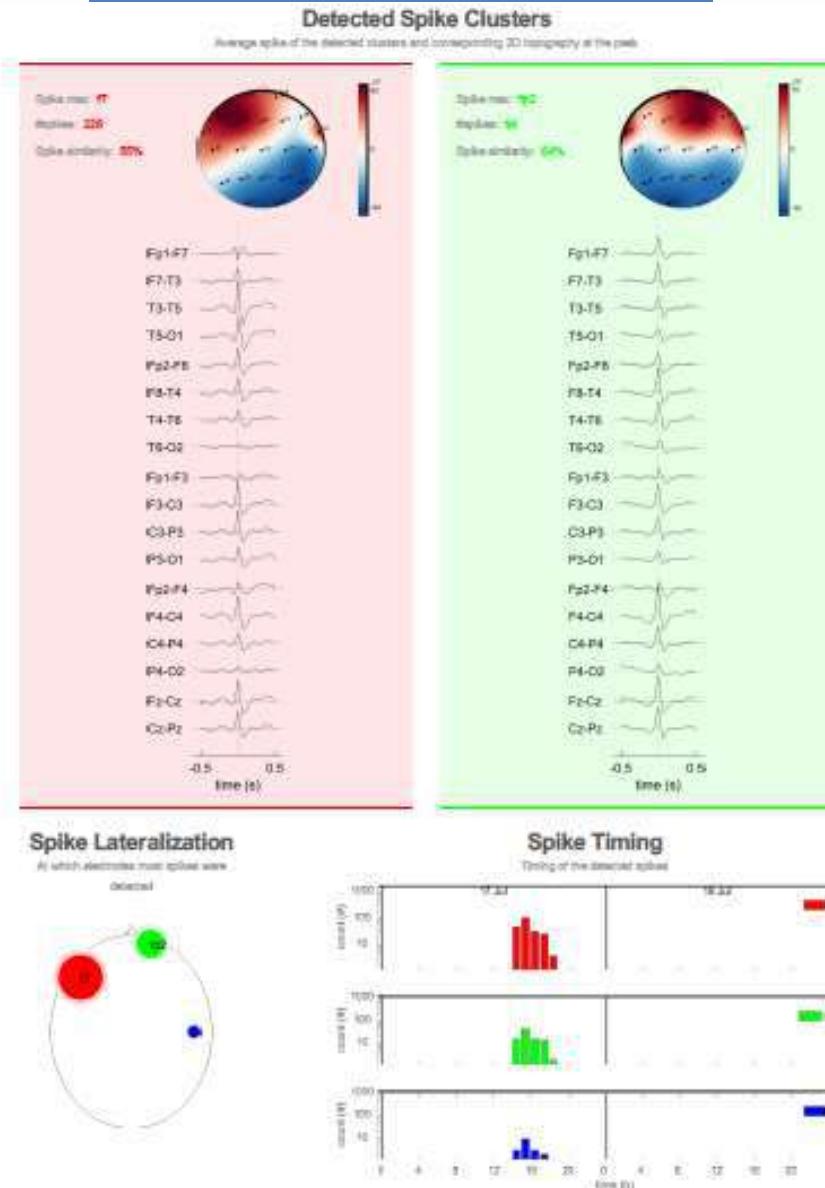


MAP07 coregistration – SPECT with MR

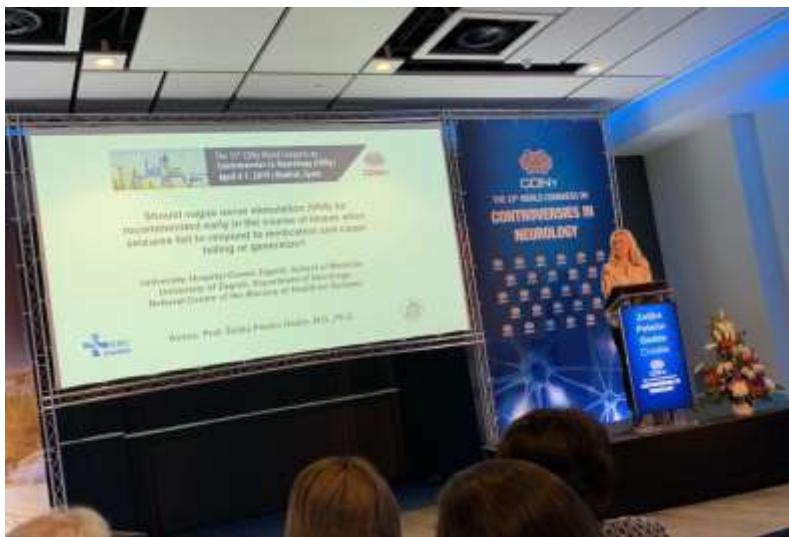




SmartBrain
08/08/2015



Automated EEG source imaging.



Coll. Antropol. 35 (2011) 3: 777-777
Original scientific paper

Vagus Nerve Stimulation in the Treatment of Patients with Pharmacoresistant Epilepsy: Our Experiences

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ABSTRACT

Vagus nerve stimulation (VNS) for the treatment of refractory partial epileptic seizures with or without generalisation in patients older than 12 years was approved in Europe in 1994 and in the United States in 2001. We studied the efficacy of VNS in patients with pharmacoresistant epilepsy hospitalized in the Neurology Department Hospital Centre Zagreb. From 1997 to 2001 we have implanted VNS in 11 patients with pharmacoresistant epilepsy, who were magnetic resonance imaging (MRI) negative and from May 2007 to May 2009 in 11 pharmacoresistant epilepsy, 9 of them were MRI positive, and were inoperable due to localisation of the pathologic changes (ganglioglioma, hamartoma, various types of cortical dysplasia, porencephalic cysts), 2 were in the group of MRI negative patients 1 patient had complex partial seizures (CPS), 6 patients had CPS with generalisation, 2 patients had primary generalized epilepsy (PGE) including myoclonic, absence, atonic and tonic seizures, one patient had PGE and CPS, and 3 patients had Lennox-Gastaut syndrome (LGS). In the group of patients one patient had elementary partial seizures (EPS) and CPS, two patients had EPS and CPS with generalisation, one patient had CPS, 3 patients had CPS with secondary generalisation, and 2 patients had secondary generalisation as well as atonic seizures. After continuous follow-up of 11 MRI negative patients 2 years and 2 MRI negative patients during one year there was decrease in mean-seizure frequency of 51.67%, 4 years follow-up of 9 MRI positive patients during 2 years there was decrease in mean-seizure frequency of 61.9%. Frequent side effects were hoarseness, throat pain and cough in the «on-phase» of the VNS, but they were mild. We can conclude that VNS was effective mode of therapy in our group of patients with pharmacoresistant epilepsy.

Key words: epilepsy, pharmacoresistance, neurosurgical treatment, vagus nerve stimulator



Case Report

Vagus nerve stimulation in Lafora body disease[☆]

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ABSTRACT

Introduction: Lafora body disease (LBD) is a rare autosomal recessive dementia and frequent nocturnal seizures in adults (GTCSE). It belongs to the group of progressive myoclonic diseases with great clinical and genetic differences, as pharmacoresistant epilepsy and progressive myoclonic seizures. We reported cases of the update of 1985 in Pöhl et al. with Engel and Haas (Möller et al., 2009).
Case presentation: A 19-year-old male patient had general epilepsy (GE), progressive cerebellar and extrapyramidal pharmacoresistance. We confirmed the diagnosis of LBD. One year follow-up period, there was a complete reduction of moderate regression of cerebellar symptomatology.
Conclusion: To our knowledge, this is the first reported case of LBD with progressive cerebellar and extrapyramidal pharmacoresistance. We confirmed the diagnosis of LBD. One year follow-up period, there was a complete reduction of moderate regression of cerebellar symptomatology.

Conclusions: To our knowledge, this is the first reported case of LBD with progressive cerebellar and extrapyramidal pharmacoresistance. We confirmed the diagnosis of LBD. One year follow-up period, there was a complete reduction of moderate regression of cerebellar symptomatology.

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Seizure outcome and use of antiepileptic drugs after epilepsy surgery according to histopathological diagnosis: a retrospective multicentre cohort study

Harm J Lambeirink^a, Willem M Otte^a, Ingmar Blümcke^b, Kees P J Braun^c, on behalf of the European Epilepsy Brain Bank writing group^d, study group^e, and the European Reference Network EpiCARE

Summary

Background Surgery is a widely accepted treatment option for drug-resistant focal epilepsy. A detailed analysis of longitudinal postoperative seizure outcomes and use of antiepileptic drugs for different brain lesions causing epilepsy is not available. We aimed to analyse the association between histopathology and seizure outcome and drug freedom up to 5 years after epilepsy surgery, to improve presurgical decision making and counselling.

Methods In this retrospective, multicentre, longitudinal, cohort study, patients who had epilepsy surgery between Jan 1, 2000, and Dec 31, 2012, at 37 collaborating tertiary referral centres across 18 European countries of the European Epilepsy Brain Bank consortium were assessed. We included patients of all ages with histopathology available after epilepsy surgery. Histopathological diagnoses and a minimal dataset of clinical variables were collected from existing local databases and patient records. The primary outcomes were freedom from disabling seizures (Engel class 1) and drug freedom at 1, 2, and 5 years after surgery. Proportions of individuals who were Engel class 1 and drug-free were reported for the 11 main categories of histopathological diagnosis. We analysed the association between histopathology, duration of epilepsy, and age at surgery, and the primary outcomes using random effects multivariable logistic regression to control for confounding.

Findings 9147 patients were included, of whom seizure outcomes were available for 8191 (89.5%) participants at 2 years, and for 5577 (61.0%) at 5 years. The diagnoses of low-grade epilepsy associated neuroepithelial tumour (LEAT), vascular malformation, and hippocampal sclerosis had the best seizure outcome at 2 years after surgery, with 77.5% (1027 of 1325) of patients free from disabling seizures for LEAT, 74.0% (328 of 443) for vascular malformation, and 71.5% (2108 of 2948) for hippocampal sclerosis. The worst seizure outcomes at 2 years were seen for patients with focal cortical dysplasia type I or mild malformation of cortical development (50.0%, 213 of 426 free from disabling seizures), those with malformation of cortical development-other (52.3%, 212 of 405 free from disabling seizures), and for those with no histopathological lesion (53.5%, 396 of 740 free from disabling seizures). The proportion of patients being both Engel class 1 and drug-free was 0–14% at 1 year and increased to 14–51% at 5 years. Children were more often drug-free; temporal lobe surgeries had the best seizure outcomes; and a longer duration of epilepsy was associated with reduced chance of favourable seizure outcomes and drug freedom. This effect of duration was evident for all lesions, except for hippocampal sclerosis.

Interpretation Histopathological diagnosis, age at surgery, and duration of epilepsy are important prognostic factors for outcomes of epilepsy surgery. In every patient with refractory focal epilepsy presumed to be lesional, evaluation for surgery should be considered.

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Seizure

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journal homepage: www.elsevier.com/locate/seizure

Vagal nerve stimulation is beneficial in postural orthostatic tachycardia syndrome and epilepsy

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Case Report

Vagus nerve stimulation in Lafora body disease[☆]Sanja Hajnsek^a, Zeljka Petelin Gadze^a, Fran Borovecki^b, Sibila Nankovic^a, Goran Mrak^c, Kristina Gotovac^b, Vlatko Sulentic^a, Ivana Kovacevic^a, Andreja Bujan Kovac^a^a University Hospital Center Zagreb and School of Medicine, University of Zagreb, Department of Neurology, Referral Centre for Epilepsy of the Ministry of Health of the Republic of Croatia, Krapinska 12, 10000 Zagreb, Croatia^b University Hospital Center Zagreb and School of Medicine, University of Zagreb, Department of Functional Genetics, Centre for Translational and Clinical Sciences, Soline 2, 10000 Zagreb, Croatia^c University Hospital Center Zagreb and School of Medicine, University of Zagreb Department of Neurosurgery, Krapinska 12, 10000 Zagreb, Croatia

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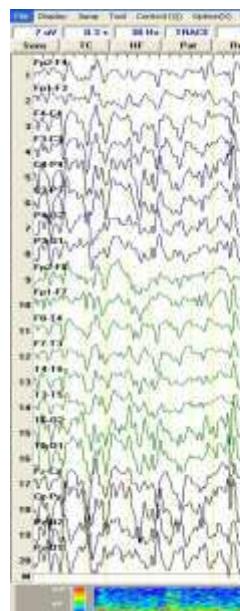
ABSTRACT

Introduction: Lafora body disease (LBD) is a rare autosomal recessive disorder characterized by progressive intractable and frequent nocturnal seizures, in addition to myoclonus and generalized tonic-clonic seizures (GTCS). It belongs to the group of progressive myoclonic epilepsies (PME), rare inherited neurodegenerative diseases with great clinical and genetic differences, as well as poor prognosis. Since these patients have a pharmacoresistant disease, an adjunctive treatment option is vagus nerve stimulation (VNS). To date, there are four reported cases of the utility of VNS in PME – in Unverricht-Lundborg disease (ULD), myoclonic epilepsy with ragged-red fibers (MERRF), Gaucher's disease, and in one case that remained unclassified.

Case presentation: A 19-year-old male patient had progressive myoclonus, GTCSs that often progressed to status epilepticus (SE), progressive cerebellar and extrapontine myopathy, and dementia, and his disease was pharmacoresistant. We confirmed the diagnosis of LBD by genetic testing. After VNS implantation, in the one-year follow-up period, there was a complete reduction of GTCS and SE, significant regression of myoclonus, and moderate regression of cerebellar symptomatology.

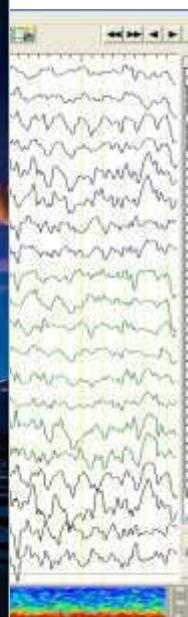
Conclusion: To our knowledge, this is the first reported case of the utility of VNS in LBD. Vagus nerve stimulation therapy may be considered a treatment option for different clinical entities of PME. Further studies with a larger number of patients are needed.

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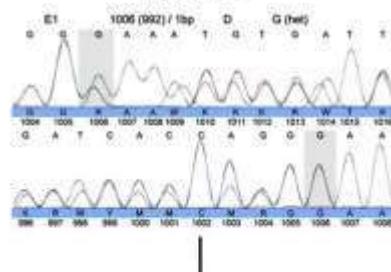
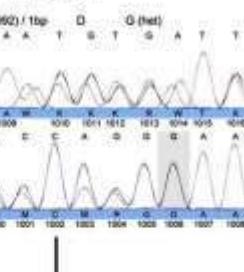
DISCOVERING HOPE

Beginning the Journey Toward Hope in Chronic Illness

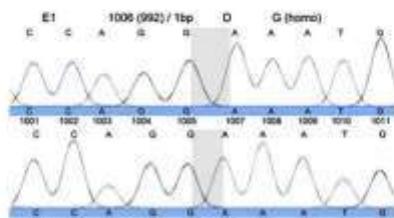


Parent 1

Parent 2



Patient



- 4/2012 – implantiran VNS (Model 103)
- 2 mA, 30-Hz signal frequency, 500-ms pulse width, 30 sec. on-time, 3 min.
- 9-godišnji period praćenja
 - potpuna redukcija generaliziranih toničko-kloničkih napadaja
 - značajna regresija mioklonizma

Dokazana homozigotna mutacija c.992delG u egzonu 1 gena EPM2B koja uzrokuje bolest Laforinih tjelešaca. Dodatne analize pokazale su u oba roditelja heterozigotnu mutaciju c.992delG.



Edukativna knjižica za bolesnike
Referentni centar Ministarstva
zdravstva RH za epilepsiju i
Hrvatska udruga za epilepsiju

Tečaj i novi sveučilišni priručnik o epilepsiji



Prof. dr. sc. Vesna Elvedi Gašparović, prof. dr. sc. Ante Čorulić i prof. dr. sc. Željka Petelin Gadžić (s lijeva na desno).

U petak, 18. listopada 2019. godine, na Medicinskom fakultetu Sveučilišta u Zagrebu, održan je poslijediplomski tečaj stalnog medicinskog usavršavanja I. kategorije "Novosti u dijagnostici i liječenju epilepsije u žena", namijenjen specijalistima neurolozima, neuropsijatrima, neonatologima, ginekologima, liječnicima obiteljske medicine, kao i specijalizantima navedenih struka. Voditeljice tečaja bile su prof. dr. sc. Željka Petelin Gadžić s Klinike za neurologiju KBC Zagreb i prof. dr. sc. Vesna Elvedi Gašparović s Klinike za ženske bolesti i porode KBC Zagreb, a stručni skup otvoren je prof. dr. sc. Ante Čorulić, ravnatelj KBC Zagreb. Bogati program tečaja s 25 predavača koncipiran je suradnjom neurologa, ginekologa, neuropsijatara, neonatologa, liječnika obiteljske medicine, specijalista laboratorijske medicine iz Hrvatske i Slovenije, s ciljem da se na



učestala neurološku bolest, a nam i-

Priručnik posvećujemo našoj dragoj profesorici Sanji Hajnšek, koja nam je prenijela entuzijazam i ljubav prema epileptologiji i uvijek nas poticala da idemo naprijed.

je promo-cija sveučilišnog priručnika "Epilepsiya - dijagnostički i terapijski pristup", koji na sveobuhvatan način na 362 stranice opisuje ovu

Mladenka Bašić, dipl. novinac
KBC Zagreb



Autori i urednici sveučilišnog priručnika "Epilepsiya - dijagnostički i terapijski pristup" - prim. mr. sc. Vlatko Šulentić, prof. dr. sc. Željka Petelin Gadžić, prim. dr. Sibila Nanković i prof. dr. sc. Zdravka Poljaković (s lijeva na desno)

EPILEPSIJA DIJAGNOSTIČKI I TERAPIJSKI PRISTUP

Urednici: Željka Petelin Gadžić, Zdravka Poljaković, Sibila Nanković, Vlatko Šulentić



THE 14th WORLD CONGRESS ON CONTROVERSIES IN NEUROLOGY (CONy)
OCTOBER 29 - NOVEMBER 1, 2020 • VIRTUAL
All times are CET (Central European)

19:50-20:30 Should surgery be offered to patients after failure of two AEDs?

Capsule: Epidemiological studies suggest that drug failure is quite likely once two agents have failed to control seizures. On the other hand, the literature contains numerous reports of response to drug therapy in patients formerly considered drug resistant. Are the ILAE guidelines supported by the evidence?

19:50-19:55 Introduction and Pre-Debate Voting

19:55-20:10 Yes: Zeljka Petelin Gadža, Croatia

20:10-20:25 No: Ettore Beghi, Italy

20:25-20:30 Rebuttals, Discussion and Post-Debate Voting





„Ljubičasti dan“ - prvi put u Hrvatskoj

• U svijetu se 26. ožujka obilježava kao „Purple Day“ („Ljubičasti dan“), Dan podrške oboljelim od epilepsije). Obilježavanje je pokrenuto 2008. godine na poticaj 9-godišnje djevojčice Cassidy Megan, njezinih roditelja i Drustva za epilepsiju Nova Scotia, Kanada.

Cassidy je prezentirala svoju bolest i probleme pred kolegama u školi te je uz majčinu podršku pokrenuta šira akcija da 26. ožujka postane Dan podrške oboljelim i da ljubičasta boja lavande postane simbol - te ona međunarodna boja epilepsije. Do danas je u akciju uključeno 15 zemalja diljem svijeta pre-

basadora. Zamisao je da se u svijetu su pristupile akciji podrške, motiviraju dobrovoljni ambasadori i promovirali „Purple Day“. Ove je godine prvi put takva akcija i u Hrvatskoj. Kanadsko Drustvo za epilepsiju prihvatio je našu aplikaciju kao ambasadori Hrvatske i namjeravaju stručnjaci epileptolozi, doc. dr. Željko Petelin i dr. Sibila Nanković.

Udruga Vlado je uključena u akciju



a riječima Cassidy Megan po svim oboljelim od epilepsije da se ne boje govorniti o svojoj bolesti. Nema vezanim uz bolest. Više „Ljubičastog dana“ u svijetu: www.purpleday.org.

Dr. Sibila Nanković
i doc. dr. Željko Petelin



Međunarodni dan epilepsije i Nacionalni dan oboljelih od epilepsije, veljača 2021.



Lejla Čarić, dr. med.



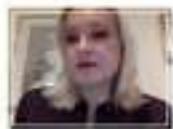
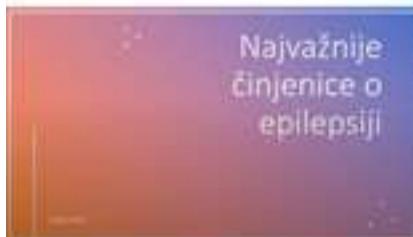
Nenad Novak, dr. med.



Ivan Lehman, dr. med.



prof. dr. sc. Boško Šparl, dr. med.



prof. dr. sc. Željka Petelin Gadžić, dr. med.



prim. Matilda Kovac Šilgorić, dr. med.



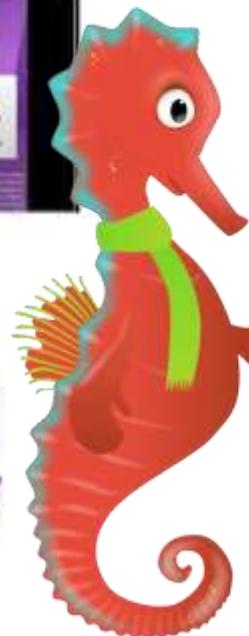
prof. dr. sc. Maia Matanović, dr. med.



prof. dr. sc. Igor Prpić, dr. med.



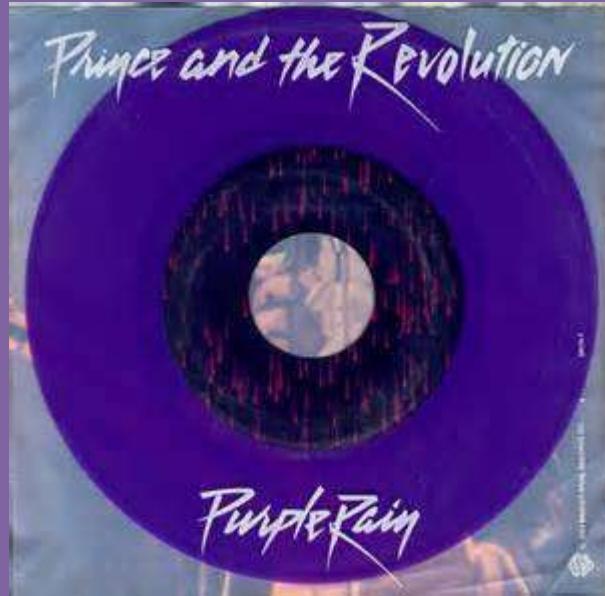
International
Epilepsy Day



KAKO POMOĆI OSOBI KOJA IMA EPILEPTIČNI NAPAD?

- 1. Ne dozvolite da vas uhvati panika!**
- 2. Mjerite vrijeme napadaja!**
- 3. Spriječite tjelesne ozljede, osobito glave!**
- 4. Osobu koja ima napadaj nakon prestanka konvulzija postavite u bočni položaj!**
- 5. Ne pokušavajte stavljati predmete između zuba!**
- 6. Ne pokušavajte napadaj prekinuti sputavanjem mišićnih kontrakcija!**
- 7. Dobro je da su u blizini samo osobe koje pružaju pomoć!**
- 8. Za vrijeme i nakon napadaja neka vaše djelovanje i način govora budu takvi da djeluju umirujuće!**
- 9. Ostanite uz osobu dok se u potpunosti ne oporavi!**
- 10. Nemojte davati osobama lijekove, hranu ili piće dok nisu u potpunosti budni!**





“I only wanna see you Laughing in the purple rain.....”

<https://youtu.be/EsRUAoUvP10>