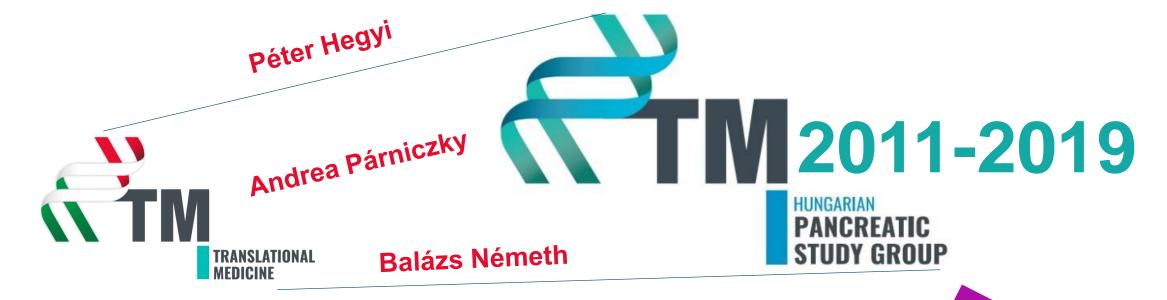


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Hungary:

Pancreatitis Working Groups and Resources



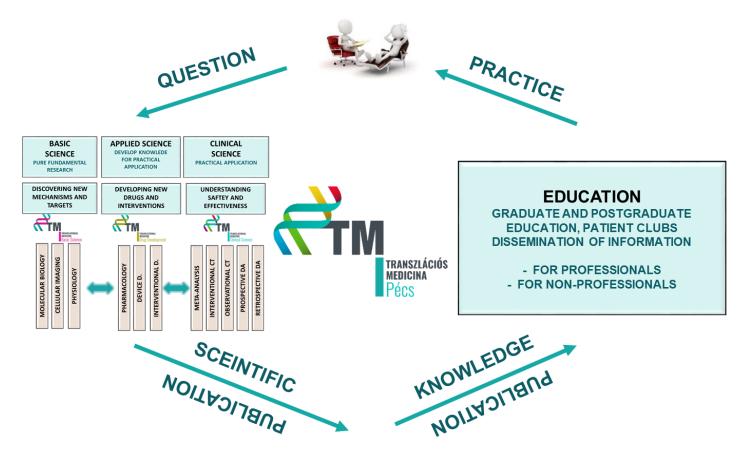




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HEALTH CARE



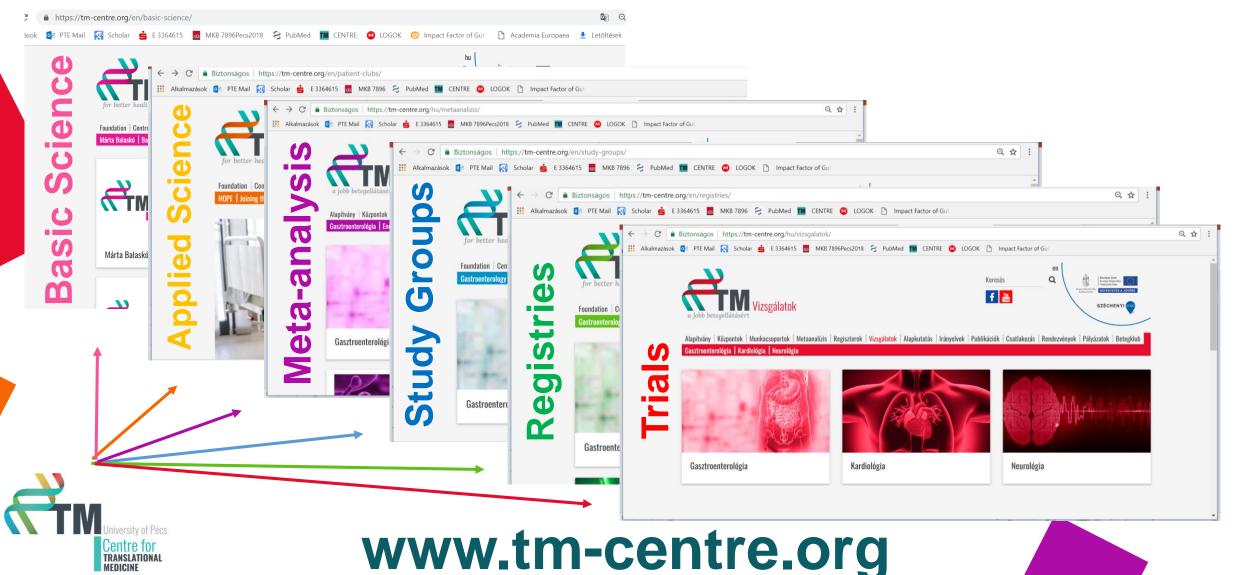
SCIENCE

KNOWLEDGE

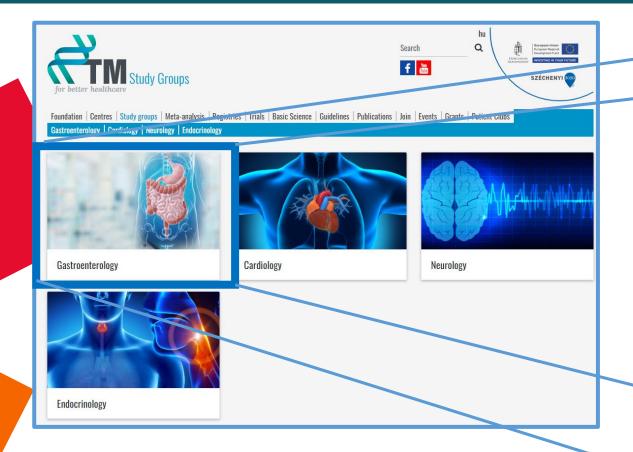
EDUCATION

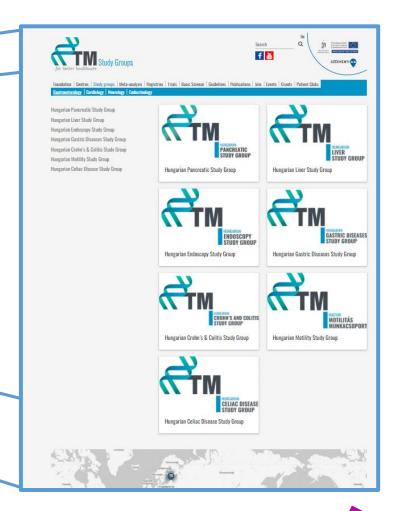
















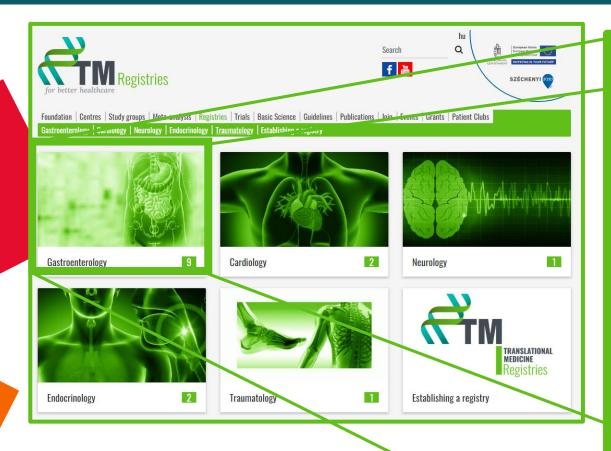
	2016 Jan	2016 Sep	2017 Feb	2017 Sep és 2018 Jan	
	ROUND I.	ROUND II.	ROUND III.	ROUND V	SUMMARY
PUBLISHED	13	14	3	4	34
ACCEPTED	0	1	1	0	2
SUBMITTED	2	3	2	8	15
IN PREPARATION	0	8	6	28	42
	15	26	12	40	93

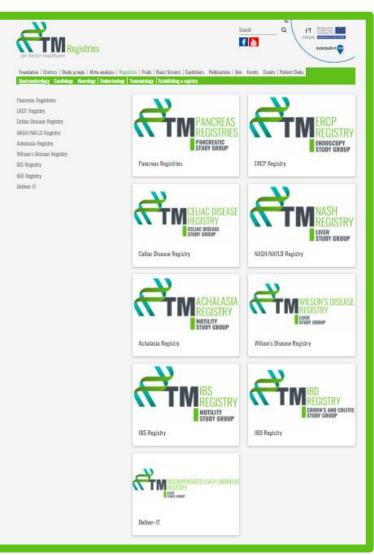
Q1	31
Q2	4
Q3	0
Q4	0
Data updated	2018.12.25



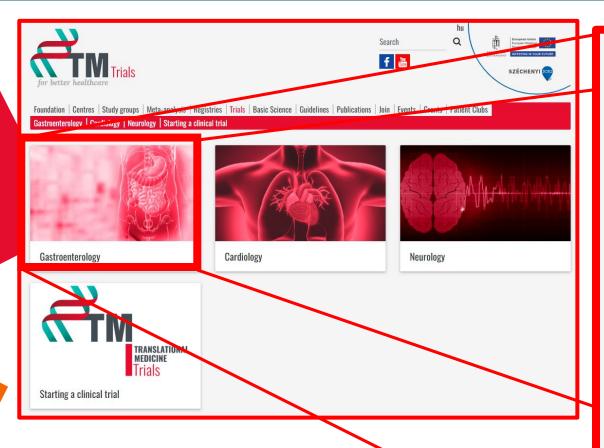


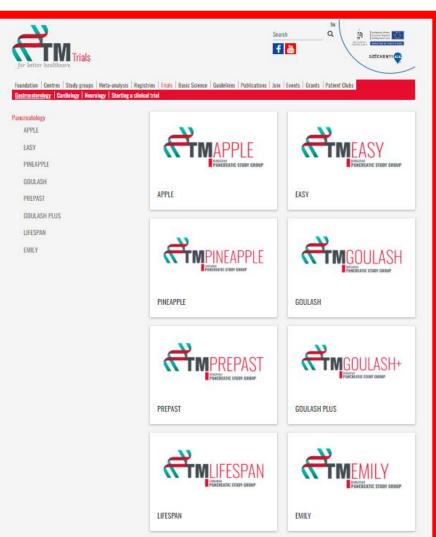












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CENTRAL INTERDISCIPLINARY UNIT





MEDICAL GROUP COORDINATORS











DATA MANAGEMENT GROUP STATISTICAL GROUP **IT GROUP**









HEALTH ECONOMICS GROUP MEDIA GROUP PATIENT CLUB COORDINATOR HR AND FINANCIAL GROUP













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ADULT (AP – CP) PANCREATITIS

Péter Hegyi



PEDIATRIC PANCREATITIS
Andrea Párniczky

GENETICS

Balázs Németh

WORKING GROUPS





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WORKING GROUPS

ADULT (AP – CP) PANCREATITIS

Péter Hegyi





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SYSTEMATIC REVIEW published: 02 April 2019 doi: 10.3389/fphys.2019.00328

frontiers
in Physiology

ORIGINAL RESEARCH published: 01 October 2018 doi: 10.3389/fphys.2018.01360

www.nature.com/scientificreports

ORIGINAL ARTICLE

OPEN

Pancreatitis-Associated Genes and Pancreatic Cancer Risk A Systematic Review and Meta-analysis

Irina Mihaela Cazacu, MD, *† Nelli Farkas, PhD, ‡ András Garami, MD, PhD, * Márta Balaskó, MD, PhD, *
Bernadett Mosdósi, MD, PhD, § Hussain Alizadeh, MD, PhD, || Zoltán Gyöngyi, MD, PhD, ¶
Zoltán Rakonczay, Jr, MD, PhD, # Éva Vigh, MD, ** Tamás Habon, MD, PhD, †† László Czopf, MD, PhD, ††
Marilena Alina Lazarescu, MD, * Bálint Erőss, MD, PhD, *
Miklós Sahin-Tóth, MD, PhD, ‡‡ and Péter Hegyi, MD, PhD, DSc(Med) § § || ||

10

Accep
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Sana Klinik

Objective: The aim of this study was to evaluate the connection between pancreatic cancer (PC) and genetic variants associated with chronic pancreatitis via systematic review and meta-analysis.

Methods: The data search was performed in 3 major databases (PubMed, Embase, and Cochrane Library). The selected studies have looked into the presence of the pancreatitis-associated genes in patients with PC and in control subjects, the outcome being the frequency of the mutations in the 2 groups. For the binary outcomes, pooled odds ratio (OR) and 95% con-

Key Words: chronic pancreatitis, CFTR, pancreatic cancer, SPINK1

(Pancreas 2018;47: 1078-1086)

P ancreatic cancer (PC) is one of the most lethal and therapeutically resistant malignancies, with a grim prognosis that is related to the late clinical presentation and the rapid progression of the disease. Despite extensive research, the etiology and pathomechanism remain



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Original Paper

Open Access Protocol

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> Protocol Open access

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BMJ Open Endoscopic sphincterotoMy for delayIng choLecystectomy in mild acute biliarY pancreatitis (EMILY study): protocol of a multicentre randomised clinical trial

> Levente Pál Kucserik, 1 Katalin Márta, 2,3 Áron Vincze, 2,4 György Lázár, 5 László Czakó, ⁶ Zsolt Szentkereszty, ⁷ Mária Papp, ⁸ Károly Palatka, ⁸ Ferenc Izbéki, ⁹ Áron Altorjay, ¹⁰ Imola Török, ¹¹ Sorin Barbu, ¹² Marcel Tantau, ¹² András Vereczkei, ¹³ Lajos Bogár, ¹⁴ Márton Dénes, ¹⁵ Imola Németh, ¹⁶ Andrea Szentesi, ^{17,18} Noémi Zádori, ² Judit Antal, ² Markus M Lerch, ¹⁹ John Neoptolemos, ²⁰ Miklós Sahin-Tóth,²¹ Ole H Petersen,²² Dezső Kelemen,²³ Péter Hegyi^{2,24}

To cite: Márta K, Szabó AN, Pécsi D, et al. High versus low energy administration in the early phase of acute pancreatitis (GOULASH trial): protocol of a multicentre randomised doubleblind clinical trial. BMJ Open 2017:7:e015874. doi:10.1136/ bmjopen-2017-015874

To cite: Kucserik LP. Márta K. Vincze Á. et al. Endoscopic sphincterotoMy for delaying choLecystectomy in mild acute biliarY pancreatitis (EMILY study): protocol of a multicentre randomised clinical trial. BMJ Open 2019;9:e025551. doi:10.1136/ bmiopen-2018-025551

Introduction According to the literature, early cholecystectomy is necessary to avoid complications related to gallstones after an initial episode of acute biliary pancreatitis (ABP). A randomised, controlled multicentre trial (the PONCHO trial) revealed that in the case of gallstone-induced pancreatitis, early cholecystectomy was safe in patients with mild gallstone pancreatitis and reduced the risk of recurrent gallstone-related

Strengths and limitations of this study

► The study is designed as a prospective, randomised-controlled trial to achieve conclusion on the highest evidence level to provide the first evidence concerning the possible benefits of sphincterotomy (ES) on timing cholecystectomy, it is (i) multinational (ii) multicentric, (iii) internationally registered and (iv) the prestudy protocol is published



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G Model YDLD-4042; No. of Pages 6

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Digestive and Liver Disease xxx (2019) xxx-xxx





ORIGINAL RESEARCH



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e First L f First L g Heim h BMKK i BMKK j Bács-l k Dr. Bu l Marku



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GOPEN ACCESS

Citation: Párniczky A, Kui B, Szentesi A, Ba Szücs Á, Mosztbacher D, et al. (2016) Pro: Multicentre, Nationwide Clinical Data from Cases of Acute Pancreatitis. PLoS ONE 11 e0165309. doi:10.1371/journal.pone.0165 Pancreatology xxx (xxxx) xxx-xxx



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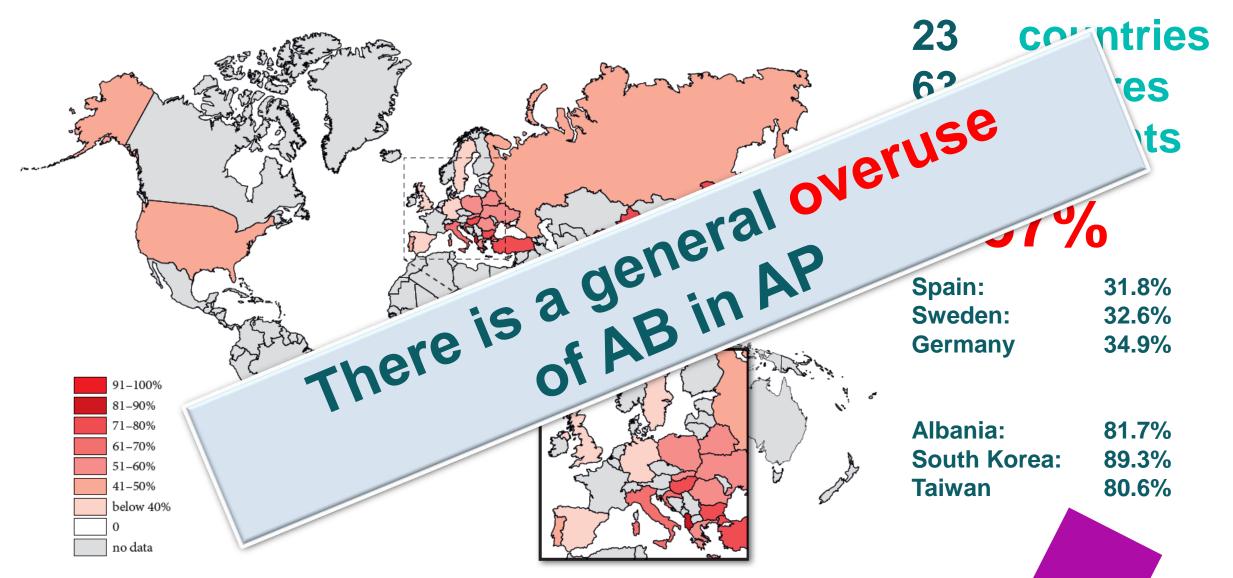
Antibiotic therapy in acute pancreatitis: From global overuse to evidence based recommendations

Andrea Párniczky a, b, 1, Tamás Lantos c, 1, Eszter Margit Tóth d, e, 1, Zsolt Szakács a, Szilárd Gódi f, Roland Hágendorn ^g. Dóra Illés ^e. Balázs Koncz ^e. Katalin Márta ^a. Alexandra Mikó ^h. Dóra Mosztbacher ^{a, i}. Balázs Csaba Németh ^e, Dániel Pécsi ^a, Anikó Szabó ^a, Ákos Szücs ^j, Péter Varjú ^a, Andrea Szentesi ^{a, e}, Erika Darvasi ^e, Bálint Erőss ^a, Ferenc Izbéki ^k, László Gaidán ^k, Adrienn Halász ^k, Áron Vincze ^f, Imre Szabó ^f, Gabriella Pár f, Judit Bajor f, Patrícia Sarlós f, József Czimmer f, József Hamvas l, Tamás Takács c, Zoltán Szepes c, László Czakó ^e, Márta Varga ^m, János Novák ^d, Barnabás Bod ⁿ, Attila Szepes ^o, János Sümegi ^p, Mária Papp ^q, Csaba Góg ^r, Imola Török ^s, Wei Huang ^t, Qing Xia ^t, Ping Xue ^u, Weiqin Li ^v, Weiwei Chen ^w, Natalia V. Shirinskaya x, Vladimir L. Poluektov y, Anna V. Shirinskaya y, Péter J. Hegyi a, z, Marian Bátovský z, Juan Armando Rodriguez-Oballe aa, Isabel Miguel Salas aa, Javier Lopez-Diaz ab, J. Enrique Dominguez-Munoz ab Xavier Molero ac, Elizabeth Pando ad, María Lourdes Ruiz-Rebollo ac, Beatriz Burgueño-Gómez ac, Yu-Ting Chang af, Ming-Chu Chang af, Ajay Sud ag, Danielle Moore ag, Robert Sutton ag, Amir Gougol ah, Georgios I. Papachristou ah Yaroslav Mykhailovych Susak ai, Illia Olehovych Tiuliukin ai, António Pedro Gomes aj, Maria Jesus Oliveira aj, David João Aparício aj, Marcel Tantau ak, Floreta Kurti al, Mila Koyacheva-Slavova am, Stephanie-Susanne Stecher an, Julia Mayerle an, Goran Poropat ao, Kshaunish Das ap, Marco Vito Marino aq, Gabriele Capurso ar, Ewa Malecka-Panas as, Hubert Zatorski as, Anita Gasiorowska at, Natalia Fabisiak at, Piotr Ceranowicz au, Beata Kuśnierz-Cabala au, Joana Rita Carvalho av, Samuel Raimundo Fernandes av, Jae Hyuck Chang aw Eun Kwang Choi ^{ax}, Jimin Han ^{ay}, Sara Bertilsson ^{az, ba}, Hanaz Jumaa ^{bb}, Gabriel Sandblom ^{bc}, Sabite Kacar ^{bd} Minas Baltatzis be Aliaksandr Vladimir Varabei bf Vizhvnis Yeshv bg Serge Chooklin bh Andriv Kozachenko bi

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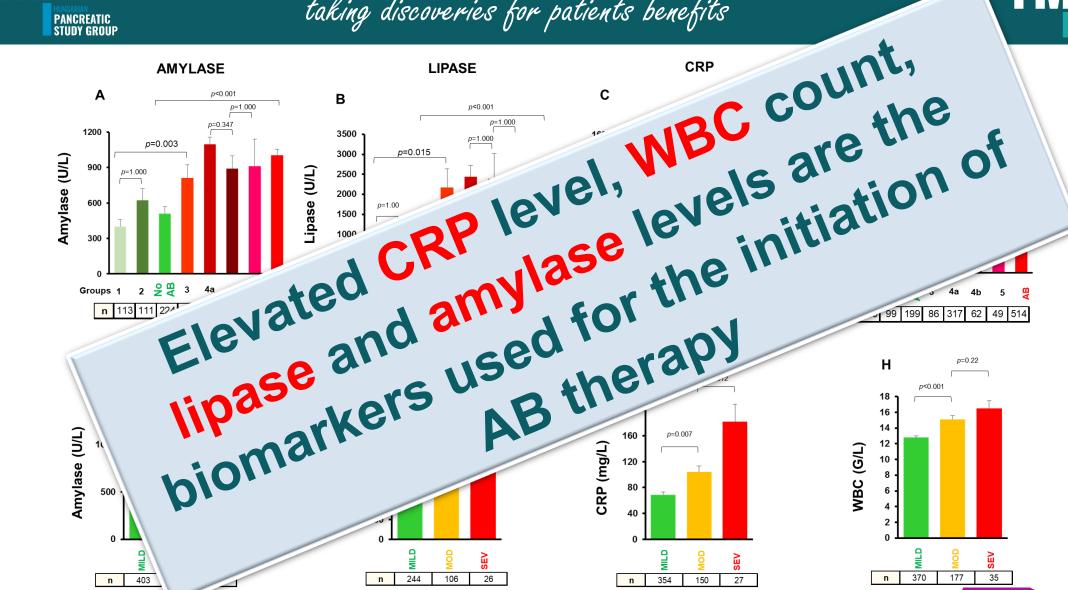
















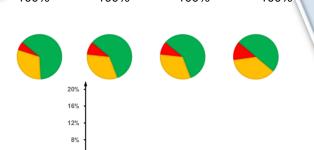
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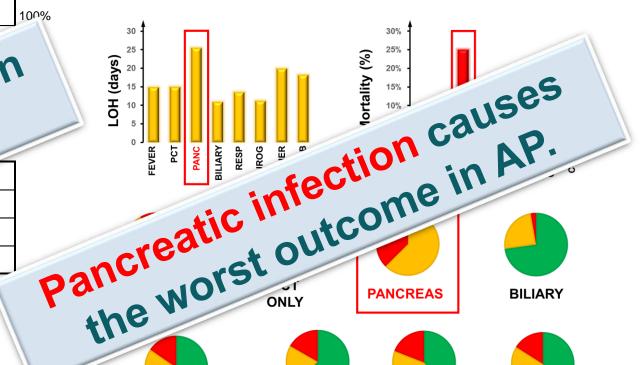
	LAB
3	8,1 ± 0,4
4a	$7,9 \pm 0,2$
4b	13,1 ± 1,3
5	16,2 ± 1.5
AB	9,6 ± 3

START OF AB THERAPY				
Day1	Day2	Day3	Day4	
72,5%	10,0%	8,3%	9,2%	100%
74,0%	11,7%	5,5%	8.80	
76,5%	6,9%	6,9%	A ir	
72,4%	10,57	start	ea.	
74,0%		eta"		

90% of AB therapy start the first 3 days of AP 54,4% 14,7% 14,7%

severity mortality





RESPIRATORY





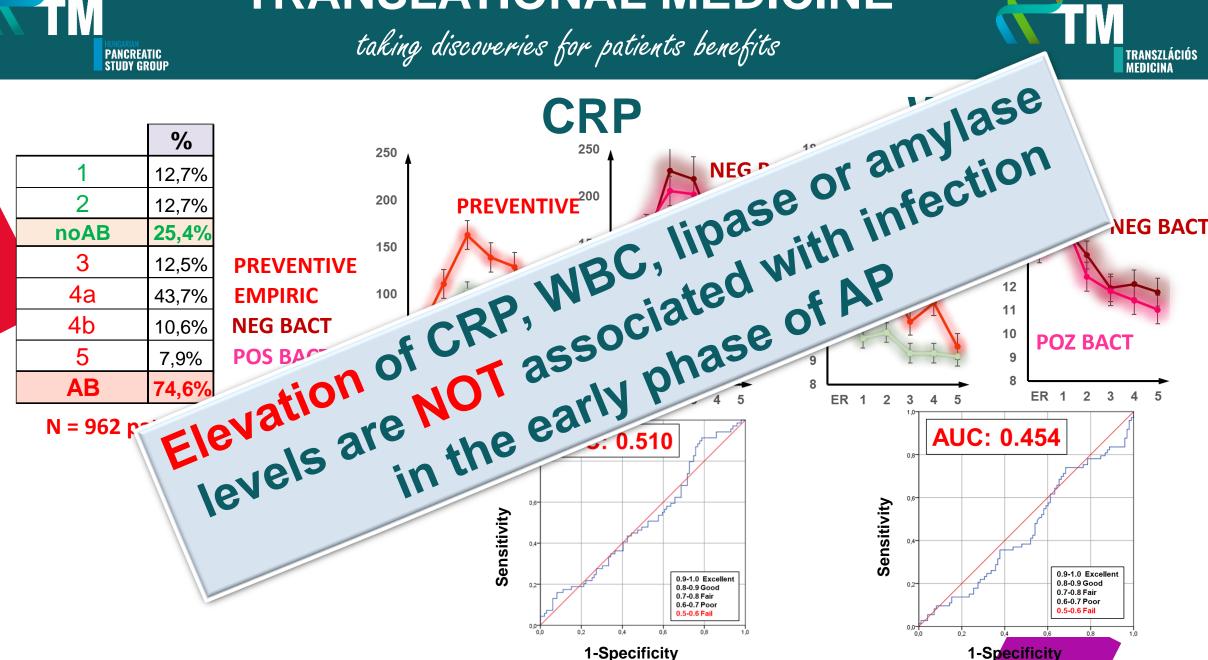


OTHER

COMBINED



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WORKING GROUPS

PEDIATRIC PANCREATITIS

Andrea Párniczky

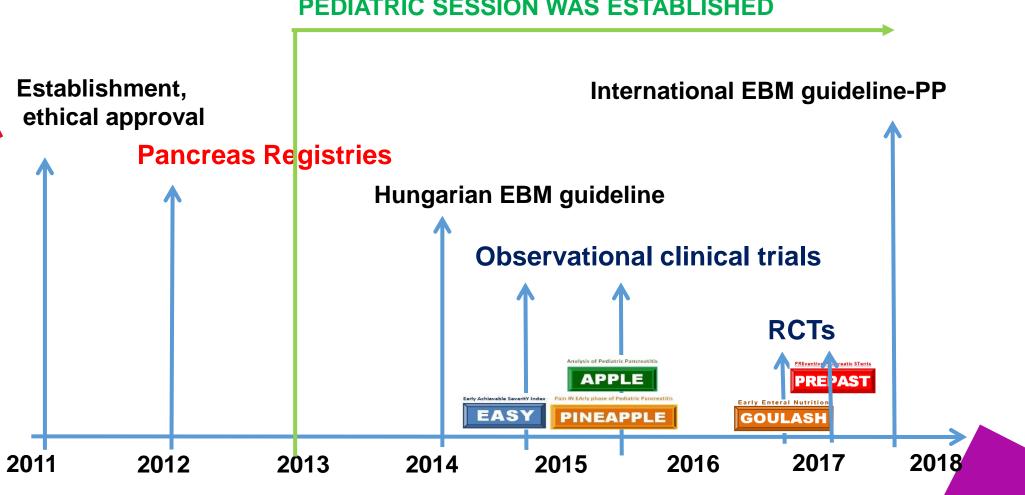




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PEDIATRIC SESSION WAS ESTABLISHED











ÖSSZEFOGLALÓ KÖZLEMÉNY

Original Paper

Digestion

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10 Depart

Pancreatology 18 (2018) 146-160

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Pancreatology

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Multina pancreatitis

Andrea Párnic Natália Lásztit

Analysi

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Group and the

^a1st Department of Group, and bDeparti dBalassa János Hosp Medical University, 5 EPC/HPSG evidence-based guidelines for the management of pediatric



Andrea Párniczky ^{a, b}, Maisam Abu-El-Haija ^c, Sohail Husain ^d, Mark Lowe ^e, Grzegorz Oracz ^f, Miklós Sahin-Tóth ^g, Flóra K. Szabó ^h, Aliye Uc ⁱ, Michael Wilschanski ^j, Heiko Witt ^k, László Czakó ^l, Tassos Grammatikopoulos ^{m, n}, Ib Christian Rasmussen ^o, Robert Sutton P. q. Péter Hegyi b, l, *

- a Heim Pál Children's Hospital, Budapest, Hungary
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- ^c Division of Gastroenterology, Hepatology and Nutrition, Cincinnati Children's Hospital Medical Center, Cincinnati, OH, USA
- d Department of Pediatrics, Children's Hospital of Pittsburgh of University of Pittsburgh Medical Center, Pittsburgh, PA, USA
- e Department of Pediatrics, Washington University School of Medicine, St. Louis, MO, USA
- f Department of Gastroenterology, Hepatology, Feeding Disorders and Pediatrics, The Children's Memorial Health Institute, Warsaw, Poland
- g Department of Molecular and Cell Biology, Center for Exocrine Disorders, Boston University Henry M. Goldman School of Dental Medicine, Boston, MA, USA
- h Division of Gastroenterology and Nutrition, Children's Hospital of Richmond, Virginia Commonwealth University, Richmond, VA, USA
- i Division of Pediatric Gastroenterology, Stead Family Department of Pediatrics, University of Iowa Carver College of Medicine, Iowa City, IA, USA
- ^j Pediatric Gastroenterology Unit, Hadassah Hebrew University Hospital, Jerusalem, Israel
- k Else Kröner-Fresenius-Zentrum für Ernährungsmedizin, Paediatric Nutritional Medicine, Technische Universität München, Freising, Germany
- ¹ First Department of Medicine, University of Szeged, Szeged, Hungary
- m Paediatric Liver, Gl & Nutrition Centre, King's College Hospital, London, United Kingdom
- ⁿ Institute of Liver Studies, Division of Transplantation Immunology and Mucosal Biology, King's College London, London, United Kingdom



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Pain IN the Early phase of Pediatric Pancreatitis

ISRCTN: 89664974

Start of trial: February, 2015

Expected closure: March, 2020

Enrolled patients:

PINEAPPLE-R: 46190

PINEAPPLE-P: 790

Number of centre: 10











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PINEAPPLE-R

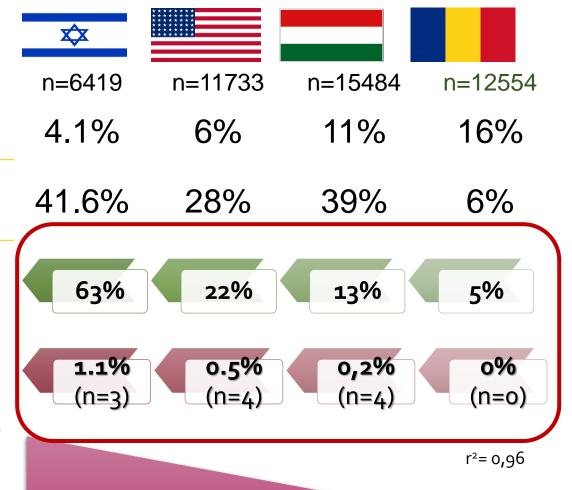
n= 46190

Abdominal pain

Abdominal imaging

Amylase/lipase

ACUTE PANCREATITIS







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PINEAPPLE-P

CURRENT PRACTICE

PPLE-P (n=790)

NEW CENTERS ARE WELCOME.

0.2%

10X

2.3%



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Analysis of Pediatric Pancreatitis

ISRCTN: 89664974

Start of trial: February, 2015

Expected closure: March, 2020

Enrolled patients:

APPLE-R: 164

APPLE-P: 79

Number of centrum: 16







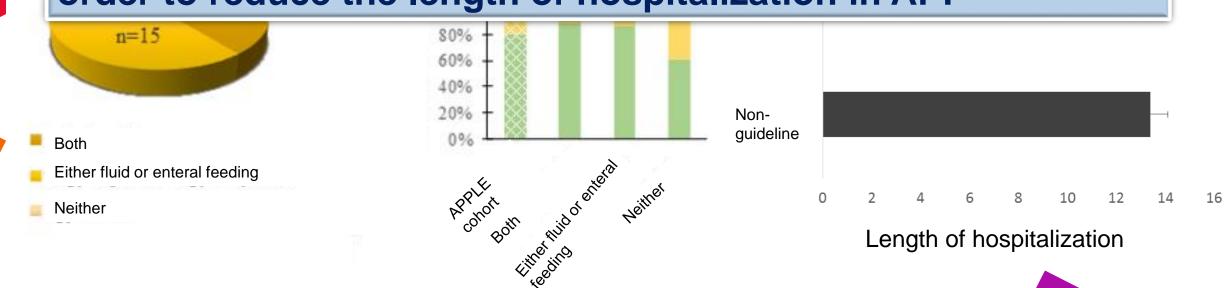




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Therapeutic strategy

Evidence-based guidelines should be followed strictly in order to reduce the length of hospitalization in APP







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	with genetic risk	all patients
	(n=59)	(n=121)
PRSS1	6.8%	2.5%

Pathogenic variants were identifications of the cases and in 34% of non-idio cases.

CFTR 25,4% 12,4% found in 32% of AP, 6 80% of CP CENTER 100 and 1 53% of idiopathic

found in 32% of AP, 65% of ARP,

■ PRSS1 ■ CTRC ■ SPINK1 ■ CPA1 ■ CFTR



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WORKING GROUPS

GENETICS

Balázs Németh



Genetic testing in Hungary for patients with pancreatitis





- 1. acute / recurrent acute / chronic pancreatitis
- 2. testing is independent from etiology
- **3. pancreatitis under 19** years of age at the time of diagnosis independent from actual age





- **1. recurrent acute pancreatitis** at least 2 acute episodes
- 2. chronic pancreatitis
- **3. pancreatitis** under 35 years of age at the time of diagnosis independent from actual age
- 4. Etiology is unknown (IDIOPATHIC)



Genetic testing in Hungary for patients with pancreatitis







MIKLÓS SAHIN-TÓTH

LETTER

Novel p. causes n heredita autosom dominan

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functionall CPA1 gen with spora

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Am J Gastroenterol. Author manuscript; available in PMC 2018 June 01.

PostScrip

Published in final edited form as:

Am J Gastroenterol. 2017 December; 112(12): 1896–1898. doi:10.1038/ajg.2017.393.

Novel PRSS1 mutation p.P17T validates pathogenic relevance of CTRC-mediated processing of the trypsinogen activation peptide in chronic pancreatitis

Balázs Csaba Németh^{1,2}, Ákos Szücs³, Péter Hegyi^{4,5,*}, and Miklós Sahin-Tóth^{1,*}

¹Center for Exocrine Disorders, Department of Molecular and Cell Biology, Boston University Henry M. Goldman School of Dental Medicine, Boston, MA 02118

²First Department of Medicine, University of Szeged, Szeged, Hungary

³First Department of Surgery, Semmelweis University, Budapest, Hungary

⁴Institute for Translational Medicine and First Department of Medicine, University of Pécs, Pécs, Hungary

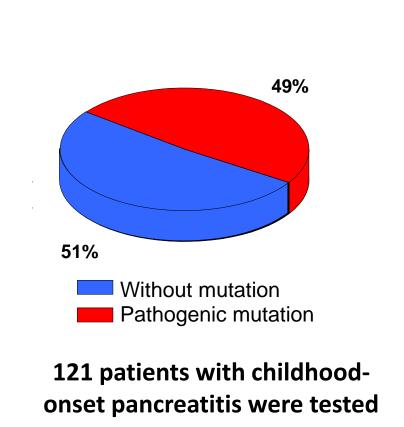
⁵MTA-SZTE Translational Gastroenterology Research Group, Szeged, Hungary

To the editor

Mutations in PRSS1 cause hereditary pancreatitis by reducing chymotrypsin C (CTRC)dependent degradation of cationic truncinggen and thereby promoting truncing





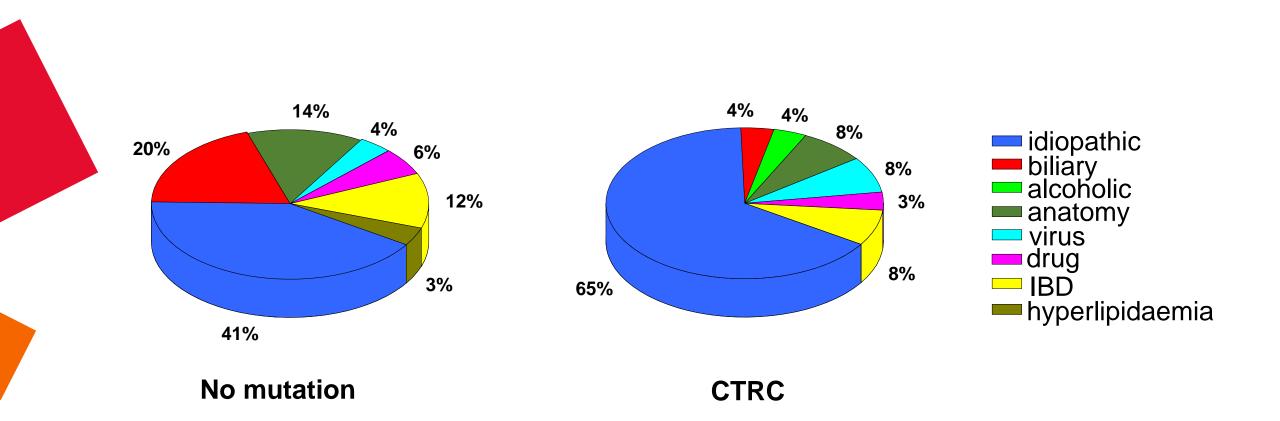


	Percentage of mutation carriers in patients
PRSS1	2.5%
CTRC	37.3%
SPINK1	9.9%
CPA1	2.5%
CFTR	12.4%



Etiology

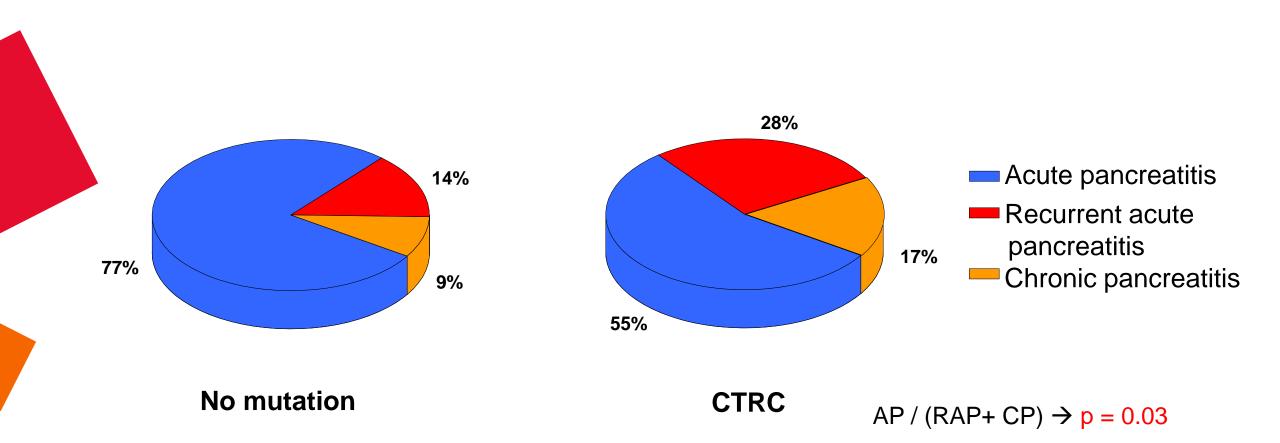






Type of pancreatitis





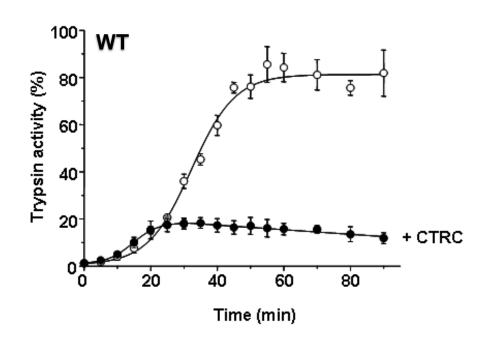


MAGYAR HASNYÁLMIR HANNYÁLMIR

PRSS1 p.P17T mutation

In a case of childhood onset chronic pancreatitis

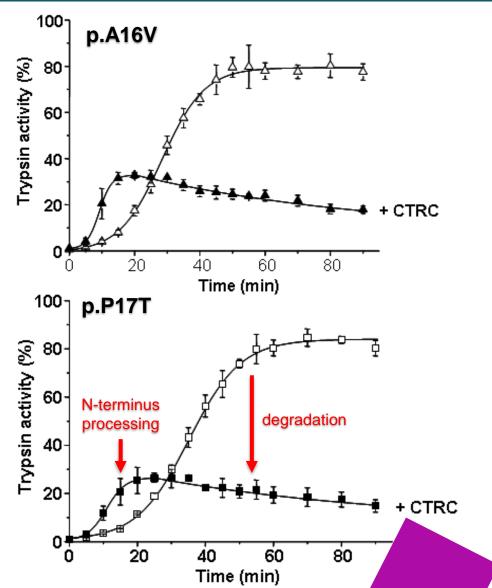




Németh BC, Szücs Á, Hegyi P, Sahin-Tóth M.

Novel PRSS1 Mutation p.P17T Validates Pathogenic Relevance of CTRC-Mediated Processing of the Trypsinogen Activation Peptide in Chronic Pancreatitis.

Am J Gastroenterol. 2017 Dec;112(12):1896-1898





EPC - BERGEN





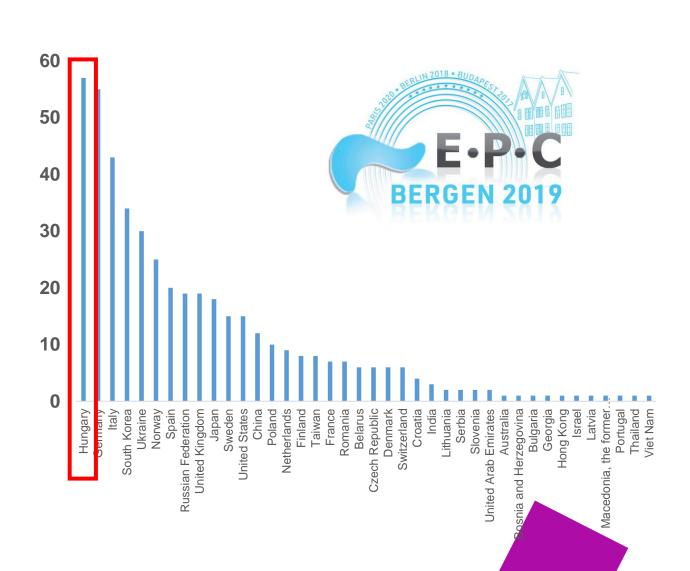


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2011: 13 abstracts



2019: 57 abstracts



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Thank you for your attention!



The Hungarian Pancreatic Study Group is committed to improving the lives of patients suffering from pancreatic diseases!









www.pancreas.hu

