Alcohol consumption and smoking dose-dependently and synergistically worsen local pancreas damage

Chronic pancreatitis (CP) is characterised by irreversible damage to the pancreas causing endocrine and exocrine dysfunction which results in decreased quality of life and reduced life expectancy.¹

Adam *et al* recently published an interesting study on a possible diagnostic tool for CP based on metabolomic profiles of patients and controls.²

Our previous international cohort analysis showed that the proportion of patients developing CP is exponentially and directly associated with the number of acute pancreatitis (AP) episodes, thus strengthening the focus on the challenging task of diagnosing CP early.^{3,4} However, in addition to diagnosing early, we should also focus on preventive interventions, before the damage becomes irreversible.

Alcohol is the main aetiological factor for CP and both alcohol consumption and smoking increase the risk for recurrence of AP and the development of CP. According to Ahmed Ali *et al*, in a follow-up study of 669 AP patients, smoking represented the dominant risk factor for recurrent AP (RAP) and a combination of alcohol consumption and smoking was the main risk factor for the progression to CP.⁵ Therefore, cessation programmes and patient education are extremely important means to intervening and lowering the recurrence of AP and the progression to CP.^{6,7} However, total cessation and abstinence often seems impossible for patients and they do not even try. Is it also possible to reduce recurrence and progression by decreasing the amount consumed?

Basic research evidence clearly suggests that alcohol and smoking amplify each other's harmful effects.^{8,9} However, large cohorts are lacking to determine whether smoking and alcohol consumption dose-dependently, mutually exacerbate the damage to the pancreas caused by each.

We have used the international cohort in the Acute Pancreatitis Registry initiated by the Hungarian Pancreatic Study Group. Data were collected from 13 countries and 30 medical centres, with 2441 cases included in the analysis. Further characteristics of the cohort and information on methods are available in online supplemental file 1.

The patient population was divided into groups according to current amounts of smoking and alcohol consumption. We found that both smoking and alcohol

Figure 1 Dose dependency in alcohol consumption and smoking in AP. (A) Light and heavy alcohol consumption and smoking groups and definitions. (B, C) Amylase and lipase levels on admission. (D) Prevalence of RAP and CP (%). (E) Proportion of local complications (%). (F, G) Amylase and lipase levels on admission. (H) Prevalence of RAP and CP (%). * $P < 0.001$, ** $p = 0.003$. Pearson's χ^2 test and the Kruskal-Wallis rank sum tests were used. AP, acute pancreatitis; CP, chronic pancreatitis; RAP, recurrent AP.

consumption are dose-dependently associated with amylase and lipase levels and with the prevalence of RAP and CP among AP patients. Alcohol consumption was also linked to a higher rate of local complications (figure 1).

Second, we examined the possible synergistic effect of these two risk factors. We arranged the cohort population into four groups based on current smoking and alcohol consumption status (figure 2). Smoking and drinking together are associated with the male sex and linked to the first AP episode 15 years earlier than non-smoking and non-drinking are. Analysing on-admission and

outcome parameters between groups, we found that amylase and lipase levels are the lowest and the proportion of moderately severe cases are the highest in the smoking–drinking group, suggesting the most pancreatic tissue damage and local complications here. The highest proportion of patients with RAP was found in both smoking groups, and the largest percentage of CP patients was observed in the smoking–drinking population, suggesting a clear synergising effect of alcohol consumption and smoking and a highlighted importance of smoking in progression.

Figure 2 Combined alcohol consumption and smoking groups in AP. (A) Groups. (B) Sex. * $P < 0.001$ in all comparisons. (C) Age. * $P < 0.001$ or $p = 0.001$ in all comparisons. (D) Severity. Moderately severe * $p = 0.003$ vs NS–ND. $P = 0.022$ vs S–ND. (E) On admission amylase level. * $P < 0.001$ vs NS–ND. ** $P < 0.003$ vs NS–ND. *** $p < 0.001$ vs NS–ND, NS–D and S–ND. (F) On-admission lipase level. * $P < 0.003$ vs NS–ND, $p < 0.006$ vs NS–D. ** $P < 0.001$ vs NS–ND and NS–D. (G) Prevalence of recurrent AP. * $P = 0.003$ vs NS–ND. ** $p = 0.04$ vs NS–D. $p < 0.001$ vs NS–ND. (H) Prevalence of chronic pancreatitis. * $P < 0.001$ vs NS–ND, NS–D. Pearson's χ^2 test and the Kruskal-Wallis rank sum tests were used. AP, acute pancreatitis; NS–D: non-smoking–drinking; NS–ND: non-smoking–non-drinking; S–D: smoking–drinking; S–ND: smoking–non-drinking.

Our analysis confirms in a clinical setting that both smoking and alcohol are dose-dependently associated with pancreatic tissue damage and the prevalence of RAP and CP. Moreover, they mutually exacerbate each other's harmful effect. In addition to the development of prognostic and therapeutic measures, further clinical trials on cessation programmes and patient education are needed. Communication to all stakeholders of the importance of at least quitting smoking or cutting the amount of smoking and alcohol consumption is crucial.

Andrea Szentesi,^{1,2} Nelli Farkas,^{1,3} Zoltán Sipos,¹ Péter Mátrai,¹ Áron Vincze,⁴ Ferenc Izbéki,⁵ Andrea Párnicsky,^{1,6} Péter Hegyi^{1,7,8} ,^{1,7,8}
Hungarian Pancreatic Study Group

¹Institute for Translational Medicine, Szentágotthai Research Centre, Medical School, University of Pécs, Pécs, Hungary

²Centre for Translational Medicine, Department of Medicine, University of Szeged, Szeged, Hungary

³Institute of Bioanalysis, Medical School, University of Pécs, Pécs, Hungary

⁴Division of Gastroenterology, First Department of Medicine, Medical School, University of Pécs, Pécs, Hungary

⁵Szent György University Teaching Hospital of Fejér County, Székesfehérvár, Hungary

⁶Heim Pál National Pediatric Institute, Budapest, Hungary

⁷Division of Pancreatic Diseases, Heart and Vascular Center, Semmelweis University, Budapest, Hungary

⁸Centre for Translational Medicine, Semmelweis University, Budapest, Hungary

Correspondence to Professor Péter Hegyi, Division of Pancreatic Diseases, Heart and Vascular Center, Semmelweis University, Budapest, Hungary; hegyi2009@gmail.com

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Collaborators Hungarian Pancreatic Study Group: Bálint Erőss, Péter Jenő Hegyi, Szilárd Váncsa, Rita Nagy, Katalin Márta, Klementina Ocskay, Márk Félix Juhász, Marcell Imrei, Mária Földi, Szabolcs Kiss, Balázs Csaba Németh, Tamás Takács, László Czakó, Szilárd Gódi, Judit Bajor, Patrícia Sarlós, László Gajdán, Mária Papp, József Hamvas, Márta Varga, Melania Macarie, Imola Török, János Novák, Artautas Mickevicius, Elena Ramirez Maldonado, Shamil Galeev, Ville Sallinen, Barnabás Bod, Ali Tüzün Ince, Tamás Nagy, Nándor Faluhelyi, Noémi Gede, Stefania Bunduc, Tamás Hussein, Mónika Lipp, Anna Németh, Orsolya Urbán, Dorottya Tarján, Simon Tóth, Dániel Pécsi, Péter Varijú, Noémi Zádori.

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ORCID iD

Péter Hegyi <http://orcid.org/0000-0003-0399-7259>

REFERENCES

- 1 Beyer G, Habtezion A, Werner J, *et al.* Chronic pancreatitis. *The Lancet* 2020;396:499–512.
- 2 Adam MG, Beyer G, Christiansen N, *et al.* Identification and validation of a multivariable prediction model based on blood plasma and serum metabolomics for the distinction of chronic pancreatitis subjects from non-pancreas disease control subjects. *Gut* 2021;70:2150–8.
- 3 Erőss B, Szentesi A, Hegyi P. Metabolic signature might be an option to identify patients with early CP. *Gut* 2021;70:2023–4.
- 4 Hegyi PJ, Soós A, Tóth E, *et al.* Evidence for diagnosis of early chronic pancreatitis after three episodes of acute pancreatitis: a cross-sectional multicentre international study with experimental animal model. *Sci Rep* 2021;11:1–14.
- 5 Ahmed Ali U, Issa Y, Hagenaars JC, *et al.* Risk of recurrent pancreatitis and progression to chronic pancreatitis after a first episode of acute pancreatitis. *Clin Gastroenterol Hepatol* 2016;14:738–46.
- 6 Nordback I, Pelli H, Lappalainen-Lehto R, *et al.* The recurrence of acute alcohol-associated pancreatitis can be reduced: a randomized controlled trial. *Gastroenterology* 2009;136:848–55.
- 7 Ocskay K, Juhász MF, Farkas N, *et al.* Recurrent acute pancreatitis prevention by the elimination of alcohol and cigarette smoking (REAPPEAR): protocol of a randomised controlled trial and a cohort study. *BMJ Open* 2022;12:e050821.
- 8 Lugea A, Gerloff A, Su H-Y, *et al.* The combination of alcohol and cigarette smoke induces endoplasmic reticulum stress and cell death in pancreatic acinar cells. *Gastroenterology* 2017;153:1674–86.
- 9 Sahin-Tóth M, Hegyi P. Smoking and drinking synergize in pancreatitis: multiple hits on multiple targets. *Gastroenterology* 2017;153:1479–81.