

## Supplementary File

### 1. Definition of ERCP volume categories and expertise levels

**ERCP volume per center** was classified as *high* (>300 procedures/year), *medium* (100–300 procedures/year), or *low* (<100 procedures/year).

**ERCP expertise** was defined as *senior* for physicians performing ERCP independently without supervision (in Slovakia, this includes formally certified endoscopists) and *junior* for those performing ERCP mostly independently but occasionally requiring guidance, including non-certified endoscopists and fellows working exclusively under supervision at the time of the survey.

**Table S1.** Overview of survey questions related to ERCP quality indicators and PEP preventive measure utilisation

Question (Number)	Close-ended versus open-ended answers
<p>5. What is your center's approach to monitoring the post-ERCP pancreatitis (PEP) rate?</p> <p><i>Diagnostic Criteria for PEP (Based on the 2012 Revised Atlanta Classification)</i>  <i>Diagnosis requires at least two of the following three criteria:</i></p> <ul style="list-style-type: none"> <li>• <i>Abdominal pain consistent with acute pancreatitis (AP).</i></li> <li>• <i>Serum amylase and/or lipase levels <math>\geq 3 \times</math> the upper limit of normal.</i></li> <li>• <i>Characteristic findings of AP on imaging (CT, MRI, or abdominal ultrasonography).</i></li> </ul>	<p>a) We monitor PEP rates using a dedicated internal registry.            b) We monitor PEP rates through periodic audits.            c) We do not monitor or collect data on PEP rates.</p>
<p>6. When tracking PEP rates, does your center include only inpatients, or both inpatients and external referrals?</p>	<p>Open-ended response.</p>
<p>7. What is your center's approach to monitoring cannulation success rates?</p>	<p>a) We routinely track cannulation rates using a dedicated internal registry that includes all procedures performed by each endoscopist.            b) We conduct periodic audits or monitor cannulation rates at least for trainees.            c) We do not monitor or collect data on cannulation rates.</p>
<p>8. How do you currently perceive — or how would you perceive, if not currently implemented — the systematic</p>	<p>a) Very positively.            b) Somewhat positively.            c) Neutral / No strong opinion.            d) Somewhat negatively.            e) Very negatively.</p>

<p>monitoring of PEP and cannulation success rates at your center?</p>	
<p>9. If you have a negative view of systematically monitoring these parameters (PEP and/or cannulation rates), or if you lack your own data, please indicate the reasons. <i>Multiple answers possible.</i></p>	<p>a) I do not want to know my or my center's data but assume it meets the required criteria.  b) I do not want to know my or my center's data because I am concerned about whether it meets the required criteria.  c) I do not consider these parameters meaningful or important for clinical practice.  d) I lack sufficient time, human resources, or a functioning methodology.  e) I lack sufficient time or human resources but would consider monitoring if a functioning methodology were introduced.  f) Other reason (<i>please specify</i>):  _____</p>
<p>10. How do you assess difficult cannulation in your routine practice?</p> <p><i>ESGE Difficult Cannulation Criteria Defined as any of the following:</i></p> <ul style="list-style-type: none"> <li>• <i>Cannulation time exceeding 5 minutes,</i></li> <li>• <i>More than 5 contacts with the papilla, or</i></li> <li>• <i>More than one unintentional pancreatic duct cannulations or opacification.</i></li> </ul>	<p>a) We record the number of contacts with the papilla.  b) We measure the time to successful cannulation.  c) We track both the number of papilla contacts and the cannulation time.  d) We count papilla contacts but estimate the cannulation time.  e) We do not record contacts or time; assessment is based on subjective judgment or experience.</p>
<p>11. What is your standard approach to biliary cannulation in patients with a native papilla?</p>	<p>a) I typically initiate cannulation using a sphincterotome.  b) I typically initiate cannulation using a standard ERCP cannula.  c) I decide on the choice of instrument on a case-by-case basis, depending on the individual patient.</p>
<p>12. What is your preferred strategy for biliary cannulation in patients with a native papilla?</p>	<p>a) I primarily use the guidewire-assisted cannulation technique.  b) I primarily use the contrast-assisted cannulation technique.  c) I do not have a preferred technique and choose the method based on the clinical situation.</p>

<p>13. How are NSAID suppositories for PEP prophylaxis administered at your center?</p>	<p>a) NSAIDs suppositories are routinely administered to all patients immediately before or after ERCP.  b) NSAID suppositories are administered to selected patients, always immediately before or after ERCP.  c) NSAID suppositories are administered to selected patients, with timing not always immediately before or after ERCP.  d) NSAID suppositories are not administered at our center.</p>
<p>14. Which NSAID suppositories do you use most frequently?</p>	<p>a) Indomethacin  b) Diclofenac  c) Indomethacin and diclofenac (used interchangeably)  d) Other (<i>please specify</i>):</p>
<p>15. If NSAID suppositories are not routinely administered to all patients immediately before or after the procedure, please describe your usual practice and explain the reasons for it.</p>	<p>Open-ended response.</p>
<p>16. Do you use aggressive hydration for PEP prophylaxis at your center?</p> <p><i>ESGE Aggressive Hydration with Lactated Ringer's Solution Protocol</i>  <i>Recommended regimen:</i>  <i>Intravenous administration of 3 mL/kg/hour during ERCP, Followed by a 20 mL/kg bolus immediately after the procedure, and</i>  <i>Continued infusion of 3 mL/kg/hour for 8 hours post-ERCP.</i></p>	<p>a) Yes, in patients with contraindications to NSAIDs (without risk of fluid overload).  b) Yes, in patients with contraindications to NSAIDs (without risk of fluid overload), in combination with other prophylactic measures (e.g., pancreatic stents).  c) Yes, in patients with contraindications to NSAIDs (without risk of fluid overload), but we follow a modified hydration protocol.  d) No, we do not use aggressive hydration.</p>
<p>17. If you use a modified hydration regimen, which type of solution do you administer, and what dosing protocol do you follow?</p>	<p>Open-ended response.</p>
<p>18. Under what circumstances do you place prophylactic pancreatic stents (PS) during biliary cannulation?  <i>Select all that apply.</i></p> <p><i>ESGE Difficult Cannulation Criteria</i>  <i>Defined as any of the following:</i></p> <ul style="list-style-type: none"> <li>• <i>Cannulation time exceeding 5 minutes,</i></li> </ul>	<p>a) PS is placed in cases of difficult cannulation with easy pancreatic stenting.  b) PS placed after first unintended PD cannulation.  e) PS is placed only rarely at our center.  f) PS is not placed at our center at all.</p>

<ul style="list-style-type: none"> <li>• More than 5 contacts with the papilla, or</li> <li>• More than one unintentional pancreatic duct (PD) cannulations or opacification.</li> </ul> <p>ESGE Easy Pancreatic Stenting (PS) Criteria Includes any of the following: Pancreatic guidewire-assisted biliary cannulation, Transpancreatic sphincterotomy, or Repeated inadvertent PD cannulation.</p>	
19. If prophylactic pancreatic stents are rarely or never used in your practice, please explain the reasons.	Open-ended response.
20. What is the typical timing at your center for checking and, if necessary, removing a prophylactic pancreatic stent after insertion?	Open-ended response.

**Table S2.** Center volume

Number of ERCPs per year	Slovakia N (%)	Czechia N (%)	Total N (%)
<100	1 (7.1)	0	1 (3.9)
100-300	9 (64.3)	3 (25)	12 (46.1)
>300	4 (28.6)	9 (75)	13 (50)

ERCP- Endoscopic retrograde cholangiopancreatography

**Table S3.** Endoscopist Workforce

Center	Slovakia- senior	Slovakia- junior	Czechia – senior	Czechia- junior
1	2	1	2	0
2	4	1	2	1
3	2	0	4	1
4	1	0	2	1
5	1	0	6	2
6	2	0	5	1
7	1	0	3	2
8	2	1	6	6
9	1	0	2	0
10	1	2	1	0
11	1	1	4	1
12	1	2	2	2
13	1	2		
14	3	1		

The median number of senior endoscopists per center was 1 (range 1–4) in Slovakia and 2.5 (range 1–6) in Czechia ( $p = 0.006$ ), while the median number of junior endoscopists per center was 1 (range 0–2) in Slovakia and 1 (range 0–6) in Czechia ( $p = 0.31$ ).

**Table S4.** Trainee involvement

Trainees involved regularly	Slovakia N (%)	Czechia N (%)	Total N (%)
Yes	9 (64.3)	10 (83.3)	19 (73.1)
No	5 (35.7)	2 (16.7)	7 (26.9)

In Czechia, 83.3% of centers reported regular trainee participation, compared with 64.3% in Slovakia ( $p = 0.37$ ).

**Table S5.** Attitude Toward Monitoring of PEP and Cannulation Rates

Answer options	Slovakia (%)	Czechia (%)	Total (%)
Very positively	13 (92.9)	3 (25)	16 (61.5)
Somewhat positively	1 (7.1)	8 (66.7)	9 (34.6)
Neutral/No strong opinion	0	1 (8.3)	1 (3.9)
Somewhat negatively	0	0	0
Very negatively	0	0	0

Slovak respondents had a significantly more positive view of this practice ( $p = 0.001$ ), with most rating it as *very positive*, whereas Czech respondents most often rated it as *somewhat positive*.

**Table S6.** Barriers to Monitoring PEP and Cannulation Rates Among Centers Lacking These Data ( $n = 12$ )

Answer options	Slovakia (%)	Czechia (%)	Total (%)
a)	1 (16,7)	1 (16,7)	2 (16,7)
b)	0	0	0
c)	0	0	0
d)	0	0	0
e)	5 (83.3)	5 (83.3)	10 (83.3)
f)	0	0	0

Answer coded (a–f) for the Question 9 correspond to those listed in Table S1.

**Table S8.** Overview of the Timing of Evaluation of Pancreatic Stent Passage and Endoscopic Extraction

Cumulative timing of evaluation (range: 24 hours–4 weeks)	Slovakia (%)	Czechia (%)	Total (%)	p-value
Never	1 (7.1)	0	1 (3.6)	1
≤ 5 days	9 (64.3)	8 (66.7)	17 (65.4)	1
≤ 7 days	11 (78.6)	11 (91.7)	22 (84.6)	0.598
> 7 days	2 (14.3)	1 (8.3)	3 (11.5)	1