

Program Application

The application must be reviewed and signed by the medical director of the endoscopy unit.

If applying for more than one unit, please provide this information for each unit on a duplicate form. This form is available for download at www.asge.org/quality/EURP.

Please check one: ☐ Nev	w Application 🗖 F	lenewal 🗖 Reins	statement Expiratio	n date, if applicable	
Name of Medical Director:					
(Please print clearly) As the medical director of this unit I hereby attest to the accuracy of all information submitted via this	Last		First		MI
application with my signature.	Medical Director S	ignature	Specialty		Date
Type of endoscopy unit: □	Office-based	☐ Ambulatory Sur	gical Center E	l Hospital-based unit	☐ HOPD only
Unit/Group Name: If your name has changed since your last application, please provide former					
Practice Manager:					
Practice Manager's Email:					
Physical Address:					
Mailing Address: if different from physical address					
City:			State: _	Zip:	
Phone:			Fax:		
Contact Information (Importa	nt! Please list your ur	it/group name exactl	y as you wish it to app	pear on your recognition ce	rtificate, if awarded.)
Indicate any institutional aff If applying for multiple unit For the purposes of the EUR Progra ownership. Please complete an app	ts regardless of at	ffiliation, total nun	nber of endoscopy	v units under your supenits, regardless of institutional	affiliation or
Indicate the examination fra	am which the unit	received energit	ation Broof of our	ront aggraditation is ro	auirod
Indicate the organization fro Accrediting Organization:	om which the unit	received accredit	ation. Proof of cur	rent accreditation is re Expiration Date:	•
Completion of the ASGE Qu Units applying to the program are end unit representative must attend the co date. (Visit					



Membership Verification

Name and membership status of endoscopists working in the unit

At least 50% of all endoscopists working in the unit must be ASGE members, with an "endoscopist working in the unit" defined as any physician, regardless of specialty, who performs 50 or more endoscopic procedures per year in the unit.

If the unit has endoscopists performing less than 50 endoscopic procedures in the unit annually, please note the following:

- The medical director of the unit must be a member of ASGE.
- While these endoscopists do not need to be listed immediately below, performance data on these endoscopists is still required to be submitted as part of the application's Quality Policy Assessment.

(Please duplicate this form to list additional endoscopists in the same unit.)

For questions regarding membership status, please contact ASGE Customer Care at 630.573.0600.

Name	ASGE m	nember?	Annual Colonoscopy Procedures	Physician Specialty GI (gastroenterologist), IM (Internal Medicine), FP (Family Practice) Surgeon or Other	E-mail address
	□ Yes	□ No			
	□ Yes	□ No			
	□ Yes	□ No			
	□ Yes	□ No			
	□ Yes	□ No			
	□ Yes	□ No			
	□ Yes	□ No			
	□ Yes	□ No			
	□ Yes	□ No			
	□ Yes	□ No			
	□ Yes	□ No			
	□ Yes	□ No			
	□ Yes	□ No			
	□ Yes	□ No			
	□ Yes	□ No			
	□ Yes				
	□ Yes	□ No			
	□ Yes	□ No			



Attestation of Guideline Adoption

The Medical Director of the endoscopy unit must attest to adopting two ASGE clinical guidelines and the CDC guideline on infection control as unit policy for all units listed on the application. By signing this form, you attest that you understand the guidelines and have adopted them as unit policy. The ASGE guidelines are linked below and published online at www.asge.org.

Name of Medical Director	Medical Director Signature	Date				
Main Practice Address: DOPTION OF ASGE CREDENTIALING GUIDELINE he "Methods of Granting Hospital Privileges to Perform Gastrointestinal Endoscopy" (Insulanz area continuo Assancement and Cuc. 2000) was repared by the Standards of Practice Committee of the American Society for Gastrointestinal Endoscopy. It represents guideline or appropriate utilization of endoscopy and is based on a critical review of the available data and expert consensus. This document is intended to provide the principles by which credentaling organizations may create policy and practical uidelines for granting gastrointestinal endoscopic privileges. Additionally, guidelines for defining continued competence, quality inprovement, and the granting of privileges for newly developed or evolving endoscopic procedures are provided. The principles et out in this document are intended to apply universally to all those who perform endoscopic procedures certify that I understand the ASGE credentialing guideline and that our unit has adopted this guideline as unit policy and will dopt any revised versions of this guideline. Name of Medical Director Medical Director Signature Date DOPTION OF ASGE REPROCESSING GUIDELINE he "Multisociety guideline on reprocessing flexible GI endoscopes: 2016 update" now the processing of the Society for Gastrointestinal Endoscopy and the Society for Geathcrare Epidemiology of America. Tordessional organizations vary in recommended practices. This document is not intended to replace these guidelines, but to omplement them, emphasizing those areas in which a broad range of professionals have reached consensus based on the valiable evidence. Certify that I understand the ASGE reprocessing guideline and that our unit has adopted this guideline as unit policy and will dopt any revised versions of this guideline. Medical Director Signature Date DOPTION OF CDC GUIDELINE FOR ISOLATION PRECAUTIONS The CDC "Guideline for Isolation Precautions: P						
(http://www.cdc.gov/hicpac/pdf/isolation/Isolation2007.pdf) is	intended for use by healthcare providers	responsible for developing, implementing and				
		esticus Asserte in Healthean Cathing - 2007"				
	-	Date				
Name of Madical Director	Madical Divastor Circustura	Dete				
		as adopted this guideline as unit policy and will				
(http://www.asge.org/uploadedFiles/Publications (public)/Pract	tice guidelines/MS guideline reprocessing GI endoscopes.pdf)	s a position statement that was published following				
ADOPTION OF ASGE REPROCESS	ING GUIDELINE					
Name of Medical Director	Medical Director Signature	Date				
		as adopted this guideline as unit policy and will				
guidelines for granting gastrointestina improvement, and the granting of priv	I endoscopic privileges. Additionally, guid ileges for newly developed or evolving e	delines for defining continued competence, quality ndoscopic procedures are provided. The principles				
prepared by the Standards of Practice	e Committee of the American Society for	Gastrointestinal Endoscopy. It represents guideline				
ADOPTION OF ASGE CREDENTIAL	ING GUIDELINE					
Main Practice Address:						
Unit/Group Name:						



Attestation of Competency

Please attest that all pertinent staff members have completed competency assessments for endoscope reprocessing, sterile medication administration (for those staff to whom it is applicable), and infection prevention in the endoscopy unit within the prior year. (Please duplicate this form, as needed, to list additional staff.)

Name of Medical Director	Medical Director Signature	Date	
Staff Name:		Date of Completion:	
Staff Name:		Date of Completion:	•
Staff Name:		Date of Completion:	
Staff Name:		Date of Completion:	
Staff Name:		Date of Completion:	
Infection Prevention			
Name of Medical Director	Medical Director Signature	Date	
Staff Name:		Date of Completion:	
Staff Name:		Date of Completion:	
Staff Name:		Date of Completion:	
Staff Name:		Date of Completion:	
Sterile Medication Administ	ration (Safe Injection Practices)		
Name of Medical Director	Medical Director Signature	Date	
otan Name.		bate of completion.	
Staff Name:		Date of Completion:	
Staff Name:		Date of Completion:	
Staff Name:		Date of Completion:	
Assessment for Endoscope	<u>Reprocessing</u>		



Quality Policy Assessment

For sample materials to assist you in completing the Quality Policy Assessment components of the application, please visit www.asge.org/quality/eurp. Your materials do not need to mirror these samples. However, many have found them useful.

Part A

Demonstrate that unit policies have been developed and adopted for continuous or intermittent assessment of the following Quality Measures, with associated performance targets for selected measures, by attaching copies of policies with dates of approval/adoption to this application. Please submit only the policies related to the following, labeling documents submitted along with this application as indicated below. Please do not staple application materials.

- Patient assessment for procedural risk before sedated procedures employing ASA score, Mallampati Score or another standardized assessment (labeled as Attachment A.1.)
- 2. Quality of preparation during colonoscopy, employing standardized criteria (labeled as Attachment A.2.)
- 3. Cecal Intubation Rate by endoscopists, during colonoscopy (labeled as Attachment A.3.)
- 4. Adenoma detection rates by endoscopist, during colonoscopy (labeled as Attachment A.4.)
- 5. Adverse event tracking, by major classes and severity, for the unit as a whole (labeled as Attachment A.5.)
- 6. **Use of Patient Satisfaction surveys** by the unit as a whole (labeled as Attachment A.6.)
 All EURP recognized units must administer a patient satisfaction survey. The policy should note the method by which your unit's patient satisfaction survey is administered. Please submit a blank copy of the survey tool currently in use.

 (ASGE's recommended survey tool can be accessed online at www.asge.org/quality, https://www.asge.org/workArea/showcontent.aspx?id=14112.)

Part B

Submit one cycle of data pertaining to the measures listed on page 6 of the application: (a) patient risk assessment documented; (b) quality of bowel preparation documented; (c) cecal intubation rate; and (d) adenoma detection rate. You will be reporting the data in aggregate and by individual endoscopist.

- In aggregate: Report the aggregate data on page 6.
- **By individual endoscopist:** Attach a supplemental document listing the performance by endoscopist. Appendix A of this application includes a suggested format. *Please de-identify the physicians, using unique identifiers (e.g., MD1, MD2).*
- In cases of suboptimal performance, if applicable, demonstrate that improvement/remediation plans have been formulated.

Remediation plans ideally include educational plan, time period anticipated for physician/staff education, details of other interventions, goal sample size, estimated time period to reach sample size, and estimated date of completion.

The data provided is confidential, considered Quality Assurance data and inadmissible. Please retain underlying data for possible future use/audit.

1.	For what type of end	doscopy unit is the award being sough	nt? (Please select one.)	
	☐ Office-based	■ Ambulatory Surgical Center	☐ Hospital-based unit	☐ HOPD only
2.	How many of the fol type?	llowing procedures did your unit do in	the last year, and how man	y physicians perform each procedure
	Colonoscopy	procedures, performed by _	endoscopists	
	EGD:	procedures, performed by _	endoscopists	
	EUS:	procedures, performed by _	endoscopists	
	ERCP:	procedures, performed by _	endoscopists	
	For ERCP, do all ph	nysicians perform > 50/year? □ Yes	□ No □ N/A	



Quality Policy Assessment continued

•	mir doc	nimum review of the last 25 or more screening/surveillance colonoscopies per endoscopist. Attach a supplemental cument listing the performance by endoscopist. (See Appendix A for a recommended format for submitting individual vsician data.)
	ls t	he data per endoscopist being submitted for the whole year, 25 consecutive cases, or otherwise?
	□ \	Year 🗖 25 cases 🗖 Other, please specify (e.g., one quarter)
	a.	Patient risk assessment documented (Number yes / Number reviewed; % Yes):/ (%) Percent ASA I:
		Percent ASA II:
		Percent ASA III:
		Percent ASA IV:
		If the assessment is not documented in \geq 95% of cases, please provide brief rationale and improvement/remediation plants
		Data demonstrating that performance by each individual endoscopist along with an improvement/remediation plan, if applicable, should be labeled Attachment B.3.a.
	b.	Quality of bowel preparation documented (Number yes / Number reviewed; % Yes):/ (%)
		Percent Adequate or better:%)
		If the preparation quality is not documented or not recorded as adequate or better in \geq 90% of cases for the entire unit, please provide brief rationale and improvement/remediation plan.
		Data demonstrating performance by each individual endoscopist along with an improvement/remediation plan, if applicable, should be labeled Attachment B.3.b.
	c.	Cecal Intubation Rate for entire unit (Number yes / Number reviewed; % Yes):/ (%)
		Range of Cecal Intubation Rate among endoscopists:% (low) to% (high)
		If the cecal intubation rate is not \geq 95% in screening and surveillance procedures for the entire unit and for each individual endoscopist, please provide brief rationale and improvement/remediation plan.
		Data demonstrating performance by each individual endoscopist along with an improvement/remediation plan, if applicable, should be labeled Attachment B.3.c.
	d.	Adenoma detection rates for unit in Screened patients > 50 Years Old
		Numerator = Number of male patients with adenomas detected =
		Denominator = Number of male patients screened =(%)
		Range of Adenoma Detection Rate for men among endoscopists:% (low) to% (high)
		Numerator = Number of female patients with adenomas detected =
		Denominator = Number of female patients screened =(%)
		Range of Adenoma Detection Rate for women among endoscopists:% (low) to% (high)
		If the adenoma detection rate for the entire unit and for each endoscopist is not \geq 30% for men and \geq 20% for women,

please provide a brief discussion and improvement/remediation plan.

Data demonstrating performance by each individual endoscopist along with an improvement/remediation plan, if

Data demonstrating performance by each individual endoscopist along with an improvement/remediation plan, if applicable, should be labeled **Attachment B.3.d.**



Quality Policy Assessment continued

Adverse events for unit as a whole (All procedures and	types	
(Number / overall procedure Number): (%)	
How many adverse events of each variety were experie	nced within the past year?	
	Outpatient Procedure	In-patient Procedure
Deaths attributable to a procedure		
Unplanned admissions within 48 hours		
Unplanned anesthesia calls to intubate (during planned moderate sedation)		
Perforations		
Bleeds requiring transfusion		
Post ERCP Pancreatitis		
What practices does your unit use to identify adverse ev Intra-procedure and post-procedure complication Change in-patient status - requirement for hotological pack 24-48 hour call back Delayed callback (> one week) post procedured other, explain:	ations recorded during visit spital admission	ut apply.)



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Quality Improvement Project Summary

Submit as an attachment [labeled **Attachment QI**] to this application a <u>summary</u> (200-300 words) of a <u>clinical</u> quality improvement project completed in your unit. Please use the *Define-Measure-Analyze-Improve-Control* format to present your project, the related outcomes and future goals. The following questions are provided as guidance; they do not need to be answered individually. **The summary provided is confidential, considered Quality Assurance data and inadmissible.**

Define your project

- What is/was the gap in quality of care?
- What were the project goals or anticipated changes you sought to achieve?

Measure your project

- What were the performance measures of interest?
- · How was the data acquired? Was it easily accessible?
- What was the baseline performance? (measurement before intervention)
- What were the targets for performance?

Analyze your project

- What local or higher-level factors contribute to defects, gaps, or variance?
- Which factors does the project address?
- What quality improvement methods and tools were utilized? (e.g., run charts, control charts, reports showing changes over time, PDSA, Lean Six Sigma)

Improve your performance

- What intervention did you pilot or implement?
- · What did repeat measurement of performance measures show?

Control summary

- What were the outcomes of the project?
- Did you achieve the project goals? If not, what did you learn? What barriers did you encounter?
- Are there any limitations to the findings? Are there additional benefits?
- Were financial benefits or cost savings realized? If so, explain.
- How will the findings be communicated?
- Are the improvements sustainable?
- Can the intervention be disseminated to the other sites as a best practice?



Application Fees and Payment Information

Application Fees

Discounts to the program apply for units meeting either or both of the following conditions. Please see the fee table below.

- A. All endoscopists in the unit are members of ASGE.
 At least 50% of unit endoscopists must be ASGE members to apply to the program.
- B. The unit participates in the GIQuIC registry. (To learn more about GIQuIC visit www.asge.org/quality/GIQuIC.)

	EURP Only		EURP + GIQuIC		
	Primary or Single Additional Units Unit		Primary or Single Unit	Additional Units	
≥ 50% Membership	\$950	\$475	\$800	\$400	
100% Membership	\$700	\$350	\$550	\$275	

Your program application will not be processed until the application fee is received. Units will have one year from the time the application fee is paid to meet all requirements. The application fee is nonrefundable.

Payment Information					
Date:					
Unit/Group Name:					
Address 1:					
Address 2:					
City:			State:	Zip:	
Phone:			Fax:		
Email:					
Method of Payment (Please check one.) 🗖 Credit Card (p	lease com	plete below) 🗖 Ch	eck payable to ASGE	
Credit Card Type:	■ Master Card	□ Visa	☐ American Exp	ress	
Card Number:				Expiration Date:	
Authorized Name on Card (please print)					
Cardholder's Signature	100100011000010000100000000000000000000				
Mail or fax completed application with p		Society for	Gastrointestinal E	ndoscopy	

American Society for Gastrointestinal Endoscopy · 3300 Woodcreek Drive · Downers Grove, IL 60515 · Phone 630.573.0600 · Fax 630.963.8332

P.O. Box 809055 Chicago, IL 60680-9055



Fax: 630.963.8332

Application Checklist

Please do not staple or bind materials.

Be sure to submit these completed materials!

□ Program application form
☐ Proof of successful and current accreditation by a recognized accrediting body (e.g., AAAHC, AAAASF, The Joint Commission, or DNV)
☐ Membership Verification form
☐ Attestation of Guideline Adoption form
☐ Attestation of Competency form
☐ Quality Policy Assessment forms (4 pages) along with attachments Please note all attachments must be labeled as instructed. Applications will be returned for labeling
☐ Quality Improvement Project Summary [labeled Attachment QI] Please note only a summary is required for submission. Complete project documentation will be returned for summarization.
☐ New member application(s) (Visit <u>www.asge.org</u> to apply today and save.)
□ Application fees

Questions regarding your application, the program or group membership?

Please contact ASGE by phone at 630.573.0600

or via email at eurp@asge.org.



Appendix A

It is suggested the unit use the following format for submitting individual physician data. Data may be submitted in other formats, such as GIQuIC reports. *Please de-identify the physicians, using unique identifiers such as MD1, MD2, etc.*

MD	# of patients w/colonoscopy	Patient Risk Assessment Documented	Cecal Intubation Rate	Quality of Bowel Prep documented as Adequate or better	Women adenoma rates	Men adenoma detection rates
MD1						
MD2						
MD3						
MD4						
MD5						
MD6						
MD7						
MD8						
MD9						
MD10						