



## ORDER FOR: Therapeutic Implantable Continuous Glucose Monitoring

Order to be completed by Ordering Health Care Provider (HCP) in its Entirety

ORDER: Implantable Gluc	ose Monitor Syster	n										
O446T - Creation of subcutaneous pocket with insertion of implantable interstitial glucose sensor, including system activation and patient training. (Initial insertion, 1 sensor included, performed and billed by health care provider, HCP)												
□ <b>0447T</b> - Removal of implantable interstitial glucose sensor from subcutaneous pocket via incision. (Billed by HCP)												
O448T - Removal of implantable interstitial glucose sensor with creation of subcutaneous pocket at different anatomic site, insertion of new implantable sensor, including system activation. (Reinsertion, 1 sensor included, performed and billed by HCP)												
A9276 - Sensor; invasive (e.g., subcutaneous), disposable, for use with interstitial continuous glucose monitoring system, 1 unit = 1 day supply												
A9277 - Transmitter;	external, for use with	n interstitial continu	uous glucose m	nonitorir	ng syste	em						
PATIENT INFORMATION												
Patient Last, First Name:	Date of								of Birth:			
Patient Address:					State:				Zip:			
City:					Phone:							
ORDERING HEALTH CARE	PROVIDER (HCP)											
HCP Last, First Name:	NPI#: _											
HCP Address:				State: Zip:								
City: Pho					none: Fax:							
MEDICAL NECESSITY*												
All shaded areas need to be completed by the HCP to document patient's need for implantable CGM and chart/EMR requirements. Patients who do not meet their insurance plan's medical coverage criteria will be asked to sign a notice of their financial responsibility in advance of the procedure. (e.g. Medicare ABN)												
Examples of Diabetes Mellitus (not all inclusive)*:  Underlying Conditions  To Specified DM												
Hypoglycemia without		·	E08.649			E11.6	49	E13.649				
Hyperglycemia		o a c o o m a	E08.65 E10.65		E11.6							
Other specified cor		mplications	E08.69					E13.6				
Without complicati		·	E08.9	E10.9			E13.		-			
	Fill in:											
1. Diagnosis ICD-10-CM	Fill in number of Insulin administrations per day:											
2. Select Insulin Administr		☐ Pump ☐ Inj			☐ Inje	ect #			ale #			
3. Fill in: HbA1c Value:		Date of HbA1c:		SMBG/Day #:			CGM:					
4. History of hypoglycemi	curring episodes o	ycemia:				☐ YES ☐ NO		□ NO				
5. Patient previously met CGM requirements by Insurer and now elects implantable CGM								\	/ES	□ №		
6. Patient requires long-term Implantable CGM (more than 72 hours) for diagnostic use:								☐ YES		□ NO*		
7. Are frequent adjustments to insulin treatment required due to glucose monitor test results?:								<u> </u>	/ES	□ NO*		
8. Patient demonstrates an understanding of technology, is capable of using the device to recognize alerts and alarms, is motivated to use the device correctly and consistently, is willing to commit to clinical visits as needed for sensor replacement, and is expected to adhere to comprehensive diabetes treatment plan.									☐ YES		□ NO*	
9. What was last date of in-person visit with treating health care provider (within last 6 months):								DATE:				
10. Routine follow-up care is expected within (fill in date, number of weeks, number of month(s), etc.):												
PRIOR AUTHORIZATION R	EVIEW											
Urgent Review  If checked, must include rationale to justify patient need for urgent request												
HEALTH CARE PROVIDER	ATTESTATION											
This document serves as an C MONITORING SYSTEM: Sensor authorized distributor. I certif- document is true, accurate ar receive their health information relevant to the patient getting	r, Smart Transmitter, all y that I am the health cand complete, to the bes on to provide benefits v g started on therapy.	associated Eversense are provider identified t of my knowledge. By	e CGM system co d in the above se y signing you agre	mponent ction and ee that ye	ts and al d I certify our Evers	associa that the sense pa	ted diabe medical tient has	etes supp necessit consente efits verifi	lies to by informed to all	oe provid nation co ow third	ed by an ntained in this parties to	

<sup>\*</sup>The medical necessity criteria used in this form was based on CMS LCDs for Implantable Continuous Glucose Monitoring (accessed 06/14/23). Other insurer requirements may vary by plan.