MEDICAL POLICY- CONTINUOUS GLUCOSE MONITORING

Continuous glucose monitoring is designed as an adjunct to and not a replacement for finger stick measurements. Ideally, measurements via the glucose monitor should be cross-checked with a finger stick measurement prior to insulin adjustments. VCHCP will approve continuous glucose monitoring equipment under the following conditions:

1) Type 1 diabetic -AND
2) Age 8 years or older (based on assessed capability of being trained to use the device in an appropriate manner) -AND
3) Device is approved for use by FDA –AND
4) Under consultation with an endocrinologist or other diabetes specialist -AND
5) The device is ordered through a VCHCP contracted DME vendor -AND
6) Best practices is being followed include a regimen of 3 or more finger sticks each day, adherence to proper diet and exercise as well as dietary and diabetic counseling and monitoring - AND
7) One of the following:
   a) Documented extreme (“brittle”) recurrent fluctuations in blood glucose measurements (i.e. symptomatic glucose levels < 50 mg/dL, >300 mg/dL)
   b) Poorly controlled blood glucose levels (unexplained hypoglycemic episodes; recurrent diabetic ketoacidosis) refractory to multiple adjustments in self-monitoring of blood glucose and insulin administration in compliant patients
   c) Frequent hospitalizations for poorly controlled diabetes
   d) Pregnant and incapable of, or poorly compliant in, self-monitoring (including one month post-partum) or are poorly controlled despite best practices (see #6 above)
   e) Documented and clinically significant Hypoglycemic Unawareness after dietary, blood glucose and behavioral adjustments and medical management (e.g. carbohydrate absorption inhibitors)
   f) Frequent nocturnal hypoglycemia despite appropriate modifications in insulin therapy and best practices (see #6 above)
   f) The patient is legally blind and blood glucose levels are poorly controlled despite best practices (special device required)

Initial authorization will be for a period of 6 months. For continued authorization of equipment, documentation of improvement in glucose control must be submitted and will be reviewed every 6 months for continued compliance and continued need.
Attachments: None
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