0617/03 Insertion of continuous glucose monitor (CGM)

Q.
Is a continuous glucose monitor considered an implanted monitoring device and coded 38285-00 [1604] Insertion of subcutaneously implanted monitoring device?

A.
A CGM system usually consists of a glucose sensor, a transmitter, and a small external monitor to view your glucose levels. A tiny electrode (glucose sensor) is inserted under the skin (usually on abdomen) using an insertion device containing a needle.

A sensor is placed into the insertion device, and with a push of a button the glucose sensor is inserted quickly and easily. The needle is then removed once the glucose sensor is in place. The glucose sensor checks glucose levels in tissue fluid and has a small adhesive patch to hold it in place for a few days and then it must be replaced with a new sensor. The sensor is connected to a transmitter that sends the information to a monitoring and display device worn on a belt or under clothing.

ACCD Coding Rule Insulin pumps (July 2015) advises that insulin pumps are not implanted in the body. In line with this advice, a continuous glucose monitor is also not classified as an implanted device and should instead be classified to 92204-00 [1866] Noninvasive diagnostic tests, measures or investigations, not elsewhere classified which is a Type C list procedure.

DECISION
As per ACCD Coding Rule Insulin pumps (July 2015), a continuous glucose monitor is not classified as an implanted device. It is classified to 92204-00 [1866] Noninvasive diagnostic tests, measures or investigations, not elsewhere classified which is a Type C list procedure.

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