

Diabetes Home Monitoring Adjudication Guideline

Rule Category:
Billing

Ref: No:
2013-BR-004

Version Control:
Version No. 5.0

Effective Date:
29/04/2018

Revision Date:
05/05/2026

Approved by:
Daman

Responsible:
Medical Standards
& Research

**Related Adjudication
Guidelines:**
Continues Glucose
Monitoring (CGM) and
External Insulin Pump

Table of Contents

1.	Abstract	3
1.1	For Members.....	3
1.2	For Medical Professionals.....	3
2.	Scope	3
3.	Adjudication Policy.....	3
3.1	Eligibility / Coverage Criteria.....	3
3.2	Requirements for Coverage	7
3.3	Non-Coverage.....	7
3.4	Payment and Coding Rules	7
4.	Denial Codes.....	8
5.	Appendices	8
5.1	References	8
5.2	Revision History	9

1. Abstract

1.1 For Members

Diabetic patients need regular blood sugar monitoring to ensure effectiveness and safety of medical management. This can be done during regular check-ups, and at home with a glucose meter as per doctor's instructions.

For patients with frequent episodes of low blood sugar, continuous monitoring might be recommended.

1.2 For Medical Professionals

Self-monitoring of blood glucose (SMBG) is an important part of glycaemic control for diabetic patients. It is mostly recommended for patients on insulin therapy and/or those on medications associated with hypoglycaemia.

Coverage of consumables and equipment for monitoring glucose control is subject to medical necessity, as well as policy terms and conditions. Allowable quantities for strips and lancets are determined by the type and severity of Diabetes.

2. Scope

This adjudication guideline clarifies the coverage criteria and limits for diabetes home monitoring consumables and equipment as part of diabetes self-management. Supplies for the administration of diabetes medications (ex. syringes and needles for insulin) are outside the scope of this guideline.

3. Adjudication Policy

3.1 Eligibility / Coverage Criteria

1. Only patients with diagnosed Diabetes are eligible for DM home monitoring consumables and/or equipment. Patients not diagnosed with Diabetes (ex. "Pre-diabetes," "Hyperglycaemia," and "Impaired Fasting Glucose," etc.) are not covered.
2. Continuous glucose monitoring (CGM) is covered under the following conditions:
 - a. Adults and children with Type 1 diabetes.
 - b. Pregnant women on insulin therapy.
 - c. Adults with Type 2 diabetes that are on long-term insulin therapy, and one of the following:
 - i. Members with recurrent severe hypoglycaemia or impaired hypoglycaemia awareness

- ii. Members having a condition or disability (including a learning disability or cognitive impairment) that causes difficulty in self-monitoring of their blood glucose by capillary blood glucose monitoring.
 - iii. They would otherwise be advised to self-measure at least 8 times a day by the treating physician.
3. Providers are required to specify the ICD10CM diagnosis code pertaining to use of insulin when billing DM-related supplies for patients on insulin therapy.
 4. Specific plan-wise coverage for diabetes home monitoring items is detailed in Table A. DM home monitoring items are not-covered for Basic plan, as stated within the corresponding General Exclusion document.

Table A. Plan-wise coverage of diabetes home monitoring consumables and equipment

Code Description	Basic	Enhanced	Thiqa
Alcohol Wipes	Not covered as per General exclusion	Covered as per SOBs	Covered
Strips, Lancets			
Platforms from home blood glucose monitor			
Spring-powered device for the lancet/ glucometer lancing device, each		Covered if with "Medical Appliances and Medical equipment" Benefit	
Home blood glucose monitor			
Sensor, transmitter, receiver; for use with interstitial continuous glucose monitoring system*			

5. Maximum allowable quantity of supplies per month and per 3 months is detailed for each type of Diabetes in Table B. Recommendations from American Diabetes Association and other international best practice references have been incorporated.
6. Self-monitoring blood glucose items (ex. Glucometer, lancets, and strips) should not be billed for the same patient with Continuous Glucose Monitoring equipment.
7. *Coverage for the sensor, transmitter, and receiver HCPCS codes will follow the Continuous Glucose Monitoring (CGM) and External Insulin Pump Guideline as per below link.

TABLE B: Maximum allowable quantity of Lancets/strips for DOH Governmental profiles [Thiqa, Basic and ABM]:

Category	Diabetes Diagnosis	Type	Criteria	Frequency	Number of Tests / Year	Max Strip Boxes QTY 50/ [A4253]	Max Lancet Boxes QTY 100/ [A4259]
Type 1 Diabetes on insulin therapy	• without CGM		Type 1 diabetes codes except Patient on CGM [check Patient history for 6 Months]	4 times per day	1,440	7 boxes/ 3 Months	4 boxes/ 3 Months
	• With CGM		Type 1 diabetes codes + Patient has a history of using CGM Codes [6 Months]	1 time per week	48	1 box/year	1 box/year
Type 2 Diabetes	• On Insulin	• on insulin more than twice daily without CGM	Type 2 diabetes codes with Z79.4 [Long term (current) use of insulin] patient under insulin Pump without CGM	4 or more times per day	1,200	6 boxes/ 3 Months	3 boxes/ 3 Months
		• on insulin therapy once or Twice Daily without CGM		2 times per day	720	4 boxes/ 3 Months	2 boxes/ 3 Months
		• on insulin with CGM	Type 2 diabetes codes + Patient has a history of using CGM Codes [6 Months]	1 time per week	48	1 box/year	1 box/year
	• On oral hypoglycaemic medications for DOH products		Type 2 diabetes codes with Z79.84 [Long term (current) use of oral hypoglycaemic drugs]	1 time per week	48	1 box/year	1 box/year
Gestational Diabetes during pregnancy (GDM)			All GDM Diabetes related codes	4 times per day	1,200	6 boxes/3 Months	3 boxes/ 3 Months

TABLE C: Maximum allowable quantity of Lancets/strips for Enhanced profiles [DOH, DHA and Riyyati]:

Unit	Box of 50	Box of 100	Type of Diabetes
50	1	0.5	Type 2 DM, Controlled, on oral hypoglycaemic medication
75	1.5	1	Type 2 DM, Uncontrolled, on oral hypoglycaemic medication
150	3	1.5	Pre-existing DM1 or 2 in pregnancy on oral hypoglycaemic medications. Type 2 DM Controlled/Uncontrolled on less frequent insulin injections/ insulin pump therapy.
300	6	3	Type 1 DM with no hypoglycaemic events or intensive insulin regimen.
375	7.5	4	Gestational diabetes mellitus initially diagnosed
450	9	4.5	Gestational diabetes mellitus on insulin Type 1 DM with hypoglycaemic events or intensive regimen/insulin pump therapy
600	12	6	Pre-existing DM1 or 2 in pregnancy on insulin

Quantities calculated as per 3 months. Processors to assume 3-months' supply unless otherwise indicated in the claim. Auditors should check history for excessive quantity.

3.2 Requirements for Coverage

- ICD, HCPCS and/or DDC codes must be coded to the highest level of specificity.
- DM-related consumables and/or equipment should only be billed by “P” providers

3.3 Non-Coverage

- DM home monitoring items are not covered for Basic and Visitor’s Plan.
- DM home monitoring equipment are not covered for plans without “Medical appliances and Medical equipment” benefit.

3.4 Payment and Coding Rules

Please apply regulator payment rules and regulations, as well as relevant coding manuals (ICD, HCPCS, etc.).

ICD10CM diagnosis code pertaining to use of insulin should be coded when billing DM-related items (strips, lancets, etc.) for patients on insulin therapy.

Use HCPCS “spring-powered device for lancet, each” or DDC “glucometer lancing device” only once in 6 months.

Billed quantity and coding should be based on the official description. For example,

- Abu Dhabi: a box of 100 strips should be reported as HCPCS “Blood glucose test or reagent strips, per 50 strips” with Quantity 2.
- Dubai and Northern Emirates: a box of 50 blood glucose test strips should be reported as DDC “Blood glucose test strips” Quantity 1 (or HCPCS “Blood glucose test or reagent strips, per 50 strips” Quantity 1)

4. Denial Codes

Code	Code Description
CLAI-012	Submission not compliant with contractual agreement between provider and payer
DUPL-002	Payment already made for same/similar service within set time frame
MNEC-004	Service is not clinically indicated based on good clinical practice, without additional supporting diagnoses/activities
NCOV-003	Service(s) is (are) not covered
PRCE-002	Service is included in another service paid

5. Appendices

5.1 References

- <https://www.england.nhs.uk/midlands/wp-content/uploads/sites/46/2019/05/3-guidelines-on-smbg-use.pdf>
- <https://www.ncbi.nlm.nih.gov/books/NBK555976/>
- <https://www.nice.org.uk/guidance/ng17/ifp/chapter/Checking-your-own-blood-glucose-and-target-levels>
- https://www.uptodate.com/contents/glucose-monitoring-in-the-ambulatory-management-of-nonpregnant-adults-with-diabetes-mellitus?search=self%20glucose%20monitoring%20&source=search_result&selectedTitle=1~150&usage_type=default&display_rank=1#H1912263947
- https://www.uptodate.com/contents/gestational-diabetes-mellitus-glucose-management-maternal-prognosis-and-follow-up?search=self%20glucose%20monitoring%20§ionRank=2&usage_type=default&anchor=H1457610429&source=machineLearning&selectedTitle=3~150&display_rank=3#H1457610429
- <https://bestpractice.bmj.com/topics/en-gb/665/treatment-algorithm>
- <https://www.doh.gov.ae/-/media/CB9099900C0D44BAAEA3B49E4DD73BED.ashx>
- Continues Glucose Monitoring (CGM) and External Insulin Pump v6.pdf

5.2 Revision History

Date	Change(s)
14/07/2013	V 1.0
15/07/2015	V 2.0 1- Disclaimer updates as per system requirement 2- Updated coverage for diabetic strips/lancets/wipes 3- Added recent references
25/03/2018	V 3.0 1- Reformatted table of the maximum allowable quantities for clarity (limits unchanged from V2.0) 2- General content update
02/05/2023	V 4.0 Medical Criteria Update
30/05/2025	V 4.1 1- Template update 2- References updated
05/05/2026	V 5.0 Update the Lancet and Strip Quantity DOH Governmental profiles [Thiqa, Basic and ABM]

Disclaimer

By accessing these Daman Adjudication Guidelines, you acknowledge that you have read and understood the terms of use set out in the disclaimer below: The information contained in this Adjudication Guideline is intended to outline the procedures of adjudication of medical claims as applied by the National Health Insurance Company – Daman PJSC (hereinafter "Daman"). The Adjudication Guideline is not intended to be comprehensive, should not be used as treatment guidelines and should only be used for the purpose of reference or guidance for adjudication procedures and shall not be construed as conclusive. Daman in no way interferes with the treatment of patient and will not bear any responsibility for treatment decisions interpreted through Daman Adjudication Guideline. Treatment of patient is and remains at all times the sole responsibility of the treating Healthcare Provider. This Adjudication Guideline does not grant any rights or impose obligations on Daman. The Adjudication Guideline and all of the information it contains are provided "as is" without warranties of any kind, whether express or implied which are hereby expressly disclaimed.

Under no circumstances will Daman be liable to any person or business entity for any direct, indirect, special, incidental, consequential, or other damages arising out of any use of, access to, or inability to use or access to, or reliance on this Adjudication Guideline including but without limitation to, any loss of profits, business interruption, or loss of programs or information, even if Daman has been specifically advised of the possibility of such damages. Daman also disclaims all liability for any material contained in other websites linked to Daman website.

This Adjudication Guideline is subject to the laws, decrees, circulars and regulations of Abu Dhabi and UAE. Any information provided herein is general and is not intended to replace or supersede any laws or regulations related to the Adjudication Guideline as enforced in the UAE issued by any governmental entity or regulatory authority, or any other written document governing the relationship between Daman and its contracting parties.

This Adjudication Guideline is developed by Daman and is the property of Daman and may not be copied, reproduced, distributed or displayed by any third party without Daman's express written consent. This Adjudication Guideline incorporates the Current Procedural Terminology (CPT®), which is a registered trademark of the American Medical Association ("AMA") and the CPT codes and descriptions belong to the AMA. Daman reserves the right to modify, alter, amend or obsolete the Adjudication Guideline at any time by providing one month prior notice.