



Guy's and St Thomas'
NHS Foundation Trust

KING'S
College
LONDON

HCL in People on Dialysis

ABCD-DTN 2025

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ABCD DTN 2025 @sugarydoc



Sufyan Hussain



@sugarydoc



A white computer keyboard is partially visible in the top left corner. A black stethoscope lies diagonally across the white surface, with its chest piece resting near the bottom left and its earpieces extending towards the top right.

Disclosures

I have delivered non-promotional continuing medical education activities, consultancy and advisory boards for
Insulet, Medtronic, Tandem, Roche, Abbott UK, Dexcom, Vertex, Menarini, Theras, Sanofi, Lilly

I am a PI on the FORWARD (VX20-880-101) trial and our institution has received unrestricted educational and research grants funding from Abbot UK and Insulet.

I have unpaid roles in the following organisations which may receive industry sponsorship:

Association of British Clinical Diabetologist, Diabetes Technology Network (vice-chair)

International Diabetes Federation Europe (board member)

European Diabetes Forum (board member)

INNODIA (Expert Input Group on beta cell replacement)

AI tools used for illustration only

I have > 30 years of personal experience of living with diabetes

Summary:

Novel therapies and AI technologies in the management of Type 1 Diabetes



HCL in Dialysis –background & theory










HCL in Dialysis – cases



HCL in Dialysis – key takeaways

Commercial Closed Loops



	Medtronic 780G	Tandem Control IQ	Cam APS	Omnipod 5
Pump				
CGM				
Target	5.5 (default), 6.1 or 6.7 mmol/L	Range 6.1-8.9 mmol/L daytime; 6.1-6.7 mmol/L overnight; 7.8-8.9 mmol/L activity	Personalised target: 4.4-11.0 mmol/L – default 5.8 mmol/L	Personalised target: 6.1mmol/L- 8.3 mmol/L
Variables	Active insulin time I:C ratio	I:C ratio Insulin sensitivity factor Basal rates	I:C ratio	I:C ratio Insulin sensitivity Factor Active insulin time
Insulin delivery	Basal insulin adjusted every 5 minutes	Basal insulin adjusted only if SG predicted to exit range	Basal insulin set to zero: extended bolus given every 10-12 minutes	Basal insulin adjusted every 5 minutes
Connectivity	Minimed Mobile and Carelink Connect App Carelink	Glooko	CAMAPS FX App – Android only Glooko	Glooko
CE license (age)	>7 years	>6 years	>1 years (Dexcom G6) ≥ 4 years (Libre 3) Pregnancy	>2 years

Nb also

Open-source (DIY) HCL systems

Griffin TP, Gallen G, Hartnell S, Crabtree T, Holloway M, Gibb FW, Lumb A, Wilmot EG, Choudhary P, Hussain S. UK's Association of British Clinical Diabetologist's Diabetes Technology Network (ABCD-DTN): Best practice guide for hybrid closed-loop therapy. *Diabet Med.* 2023 Jul;40(7):e15078. doi: 10.1111/dme.15078. Epub 2023 Apr 10.






Received: 18 January 2023 | Accepted: 28 February 2023

DOI: 10.1111/dme.15078

DIABETIC
Medicine

POSITION STATEMENT

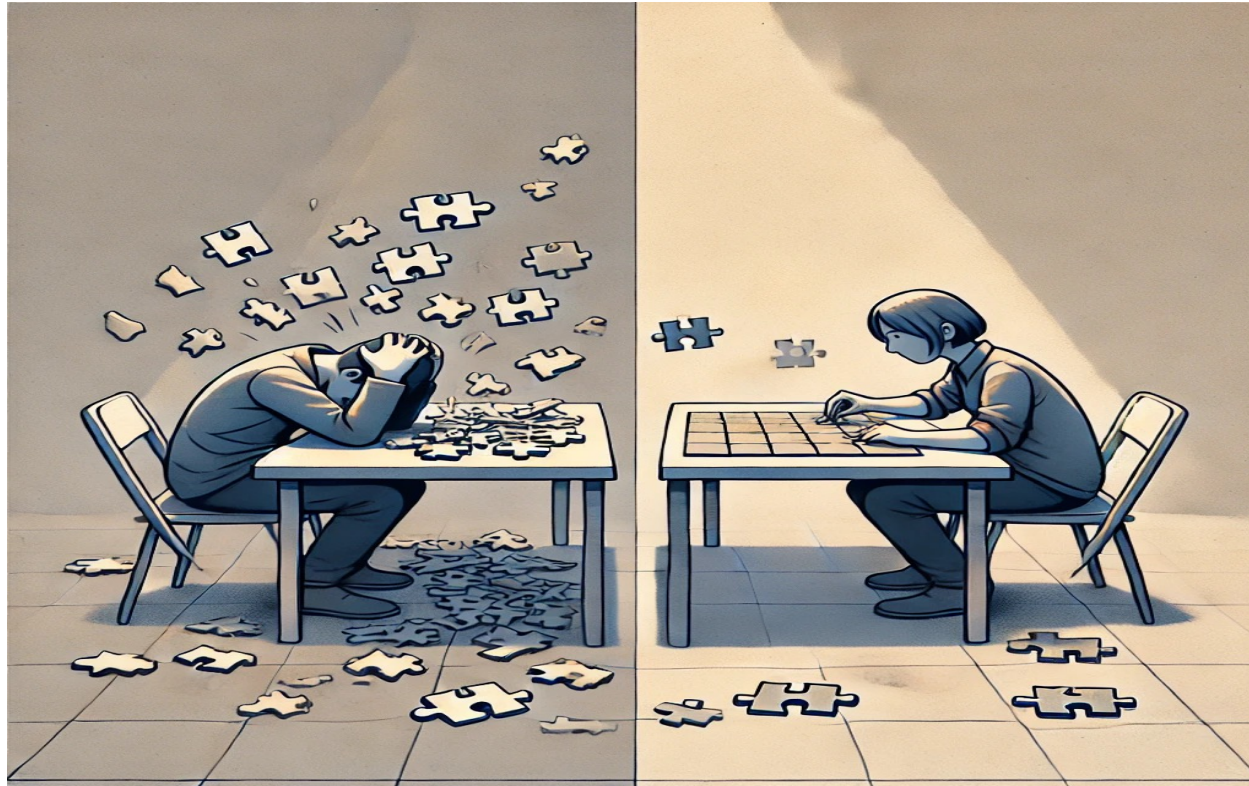
UK's Association of British Clinical Diabetologist's Diabetes Technology Network (ABCD-DTN): Best practice guide for hybrid closed-loop therapy

Tomás P. Griffin^{1,2,3,4}  | Geraldine Gallen⁵ | Sara Hartnell⁶ | Thomas Crabtree^{7,8} |
Melissa Holloway⁹ | Fraser W. Gibb¹⁰  | Alistair Lumb^{11,12}  | Emma G. Wilmot^{7,8} |
Pratik Choudhary^{1,2}  | Sufyan Hussain^{13,14} 



Griffin, TP, et al. Diabet Med. 2023; 40:e15078. doi:10.1111/dme.15078

What is complex for me may not be complex for you!



What's complex for me

Haemodialysis

Gastroparesis

Severe mental health and behavioural challenges

Extreme insulin resistance

High-dose steroid use

Visual and manual dexterity issues

Altered physiology – gastrointestinal, insulin absorption, liver metabolism

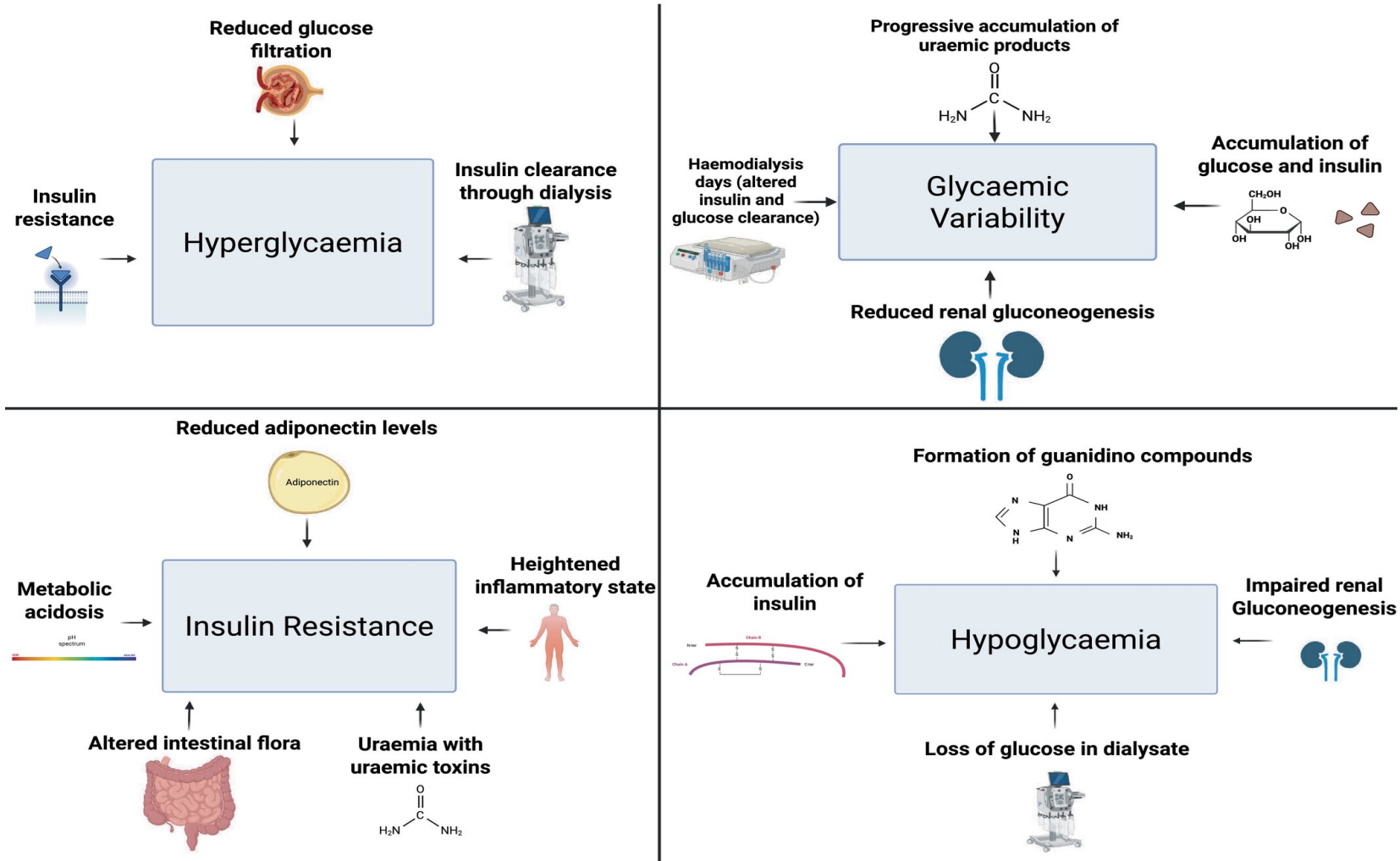
Evidence in these scenarios

- Limited
- Trials and pivotal studies
 - Majority industry supported
 - Those with extensive complications and higher levels of glycaemia often excluded
 - Bias towards more engaged and motivated

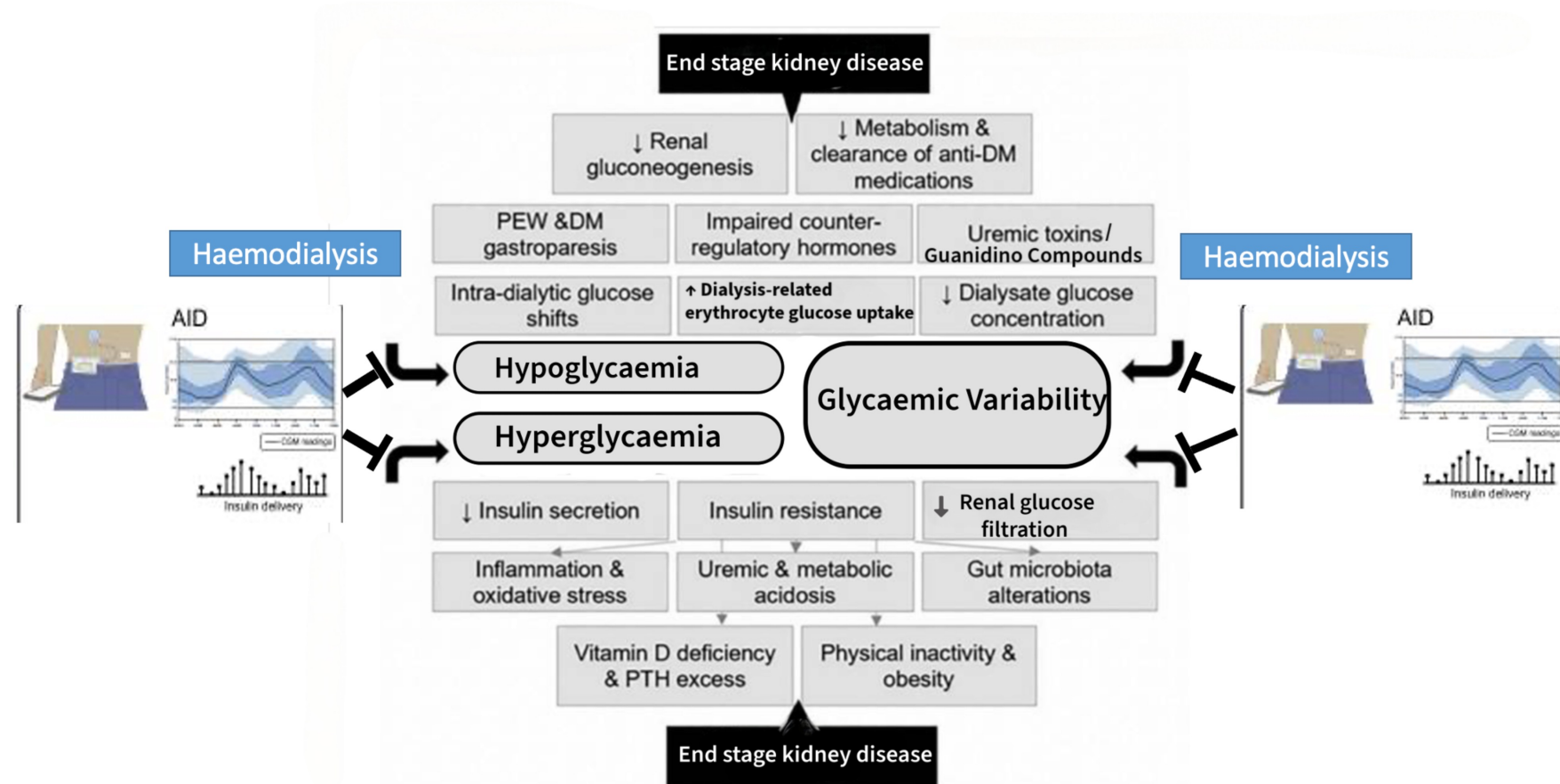
Real-world evidence

- Access and uptake of technology influenced by social determinants of health
 - Insurance driven → higher income
 - Public systems → "inverse care law"
- HCL : built on principle of "ONE SIZE FITS ALL (or most)"

Using automated insulin delivery to address the clinical challenges of glycemic management in people with type 1 diabetes and kidney failure on maintenance hemodialysis



Contributors to challenges faced by ESKD on AID



Mitigating features of AID against pathophysiological challenges of ESKD

Pathophysiological challenges in ESKD	AID mitigating feature
<ul style="list-style-type: none">• Increased risk of hypoglycaemia	<ul style="list-style-type: none">• Automatic suspension of insulin delivery upon detection of impending hypoglycaemia• Low glucose alarms helpful in nocturnal hypoglycaemia or cases of impaired hypoglycaemia awareness• Algorithms can be adjusted to prevent hypoglycaemia on dialysis days by altering glucose target or by using “activity/exercise mode”
<ul style="list-style-type: none">• Impaired insulin clearance	<ul style="list-style-type: none">• Most systems allow adjustment of active insulin time
<ul style="list-style-type: none">• Increased insulin resistance• High glucose variability	<ul style="list-style-type: none">• Automatic adjustment of basal insulin and correction bolus delivery factor at every few minutes based on CGM and pump data.• Healthcare professionals can access AID data illuminating discernible patterns in insulin requirements glucose variability, allowing for meaningful intervention thereby reducing the risk of clinical inertia due to the complexity of glycaemic profile

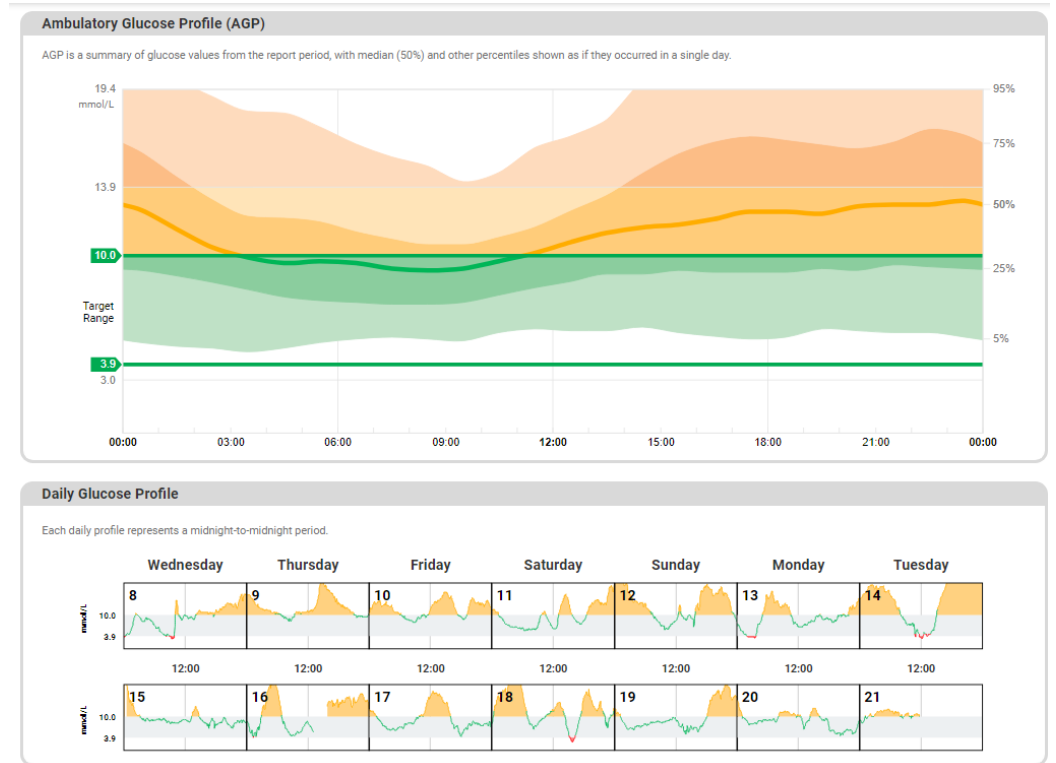
Abbreviations: ESKD=End stage kidney disease, AID= Automated insulin delivery, CGM= Continuous glucose monitor



Panagiotis Pavlou

Haemodialysis– Pre-HCL

- 35 y/o Sub-continental Asian female
- Type 1 diabetes - SPK in 2016
- Pancreas graft failed in 2017, kidney failed in 2022
- Pre-pump MDI and Dexcom G6
- TDD 16.5 units
- Pre pump data:
 - Time in range 31%
 - Time High 33% and V High 35%
 - Time below range 0%



Becky Hyslop
Thomas Johnson



Dr Janaka Karalliedde

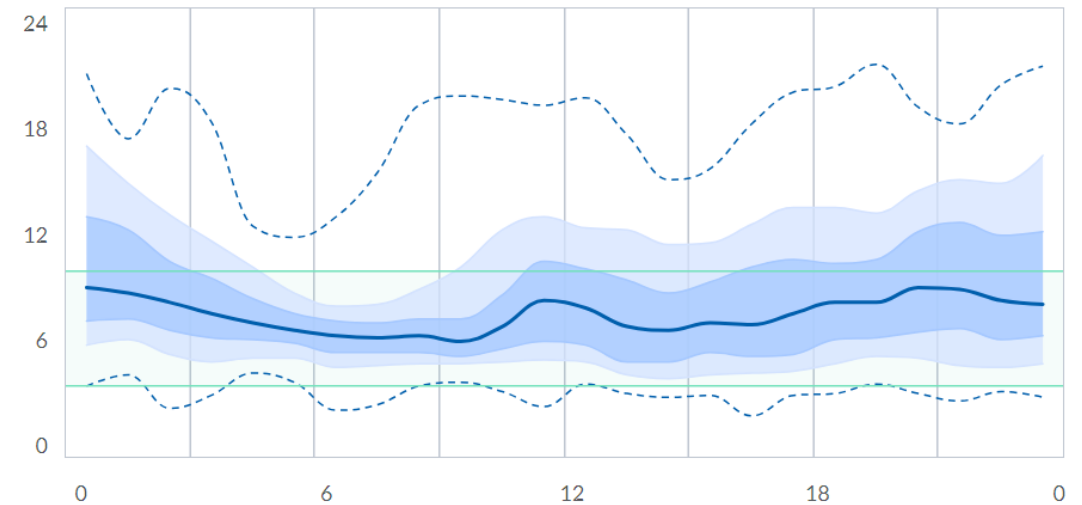
Post-HCL: CamAPS FX

- Glucose target gradually reduced to 6.5 mmol/L (117 mg/dl)
- Time in range 59%, Low 2%, High 22%, V High 17%
- GMI 7.1%
- Reduced glucose variability
 - still 39.5%
- **More hypoglycaemia**
- TDD 12.9 units

AGP

Glucose (mmol/L)

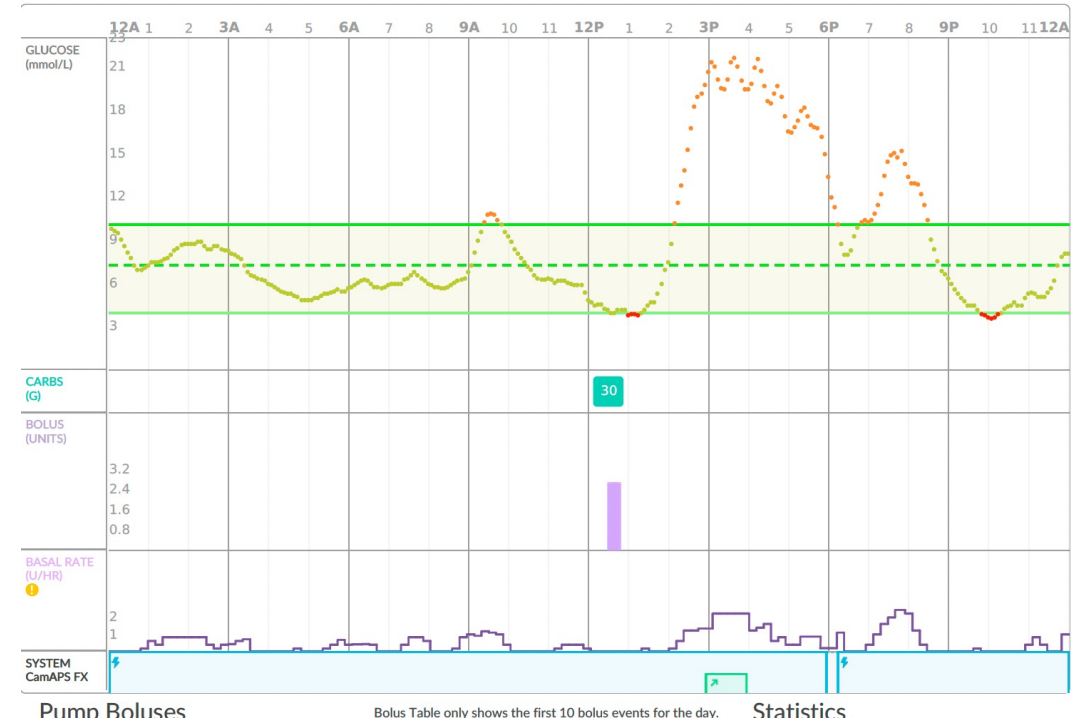
[What is AGP?](#)



Target Range (3.9 - 10 mmol/L) 25 - 75% Lowest - Highest
Median 10 - 90%

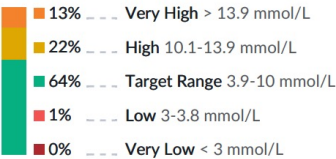
Post-HCL: CamAPS FX

- Algorithm tended to over-correct highs (delayed lows)
- Cautious use of Boost function – risk of delayed hypos
- Fear of hypos/ bolus- omitted/ delayed bolus
- Connectivity issues



- Consider using activity mode (ease off) prior to/during dialysis sessions
- Reduce bolus prior to dialysis
- Increased target?

Glucose - Time In Range

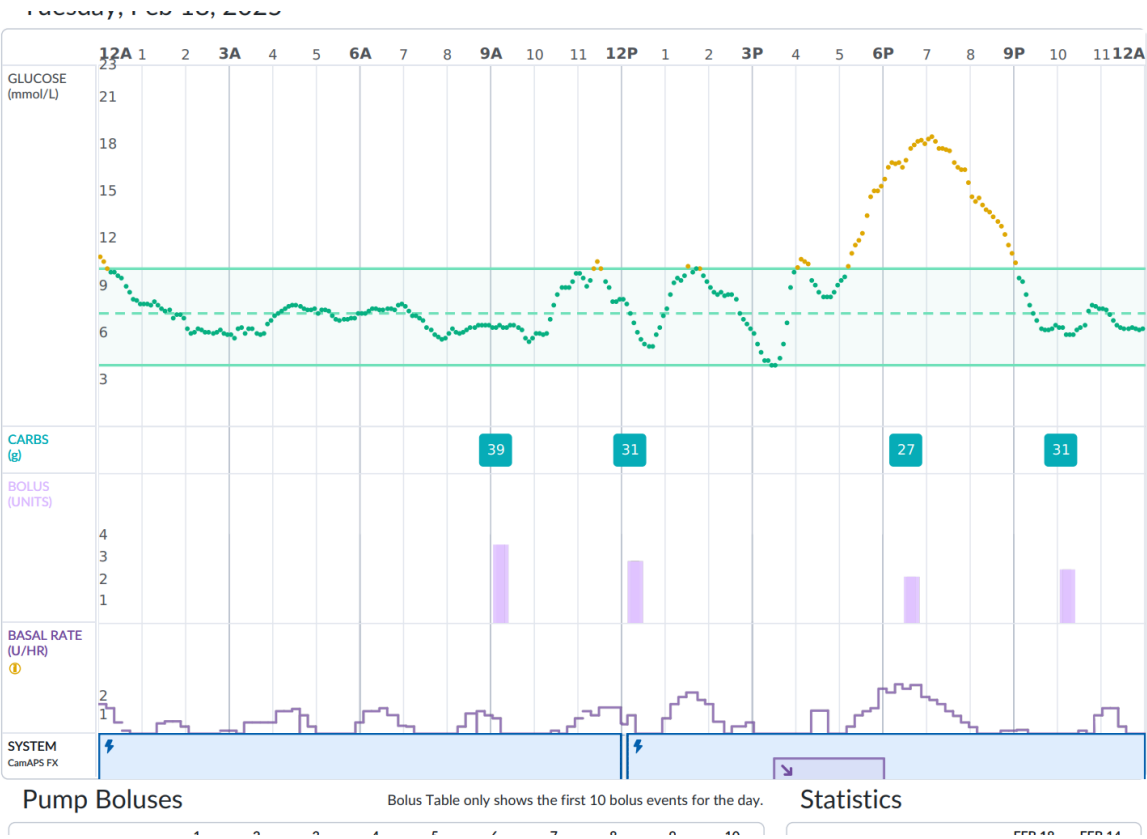
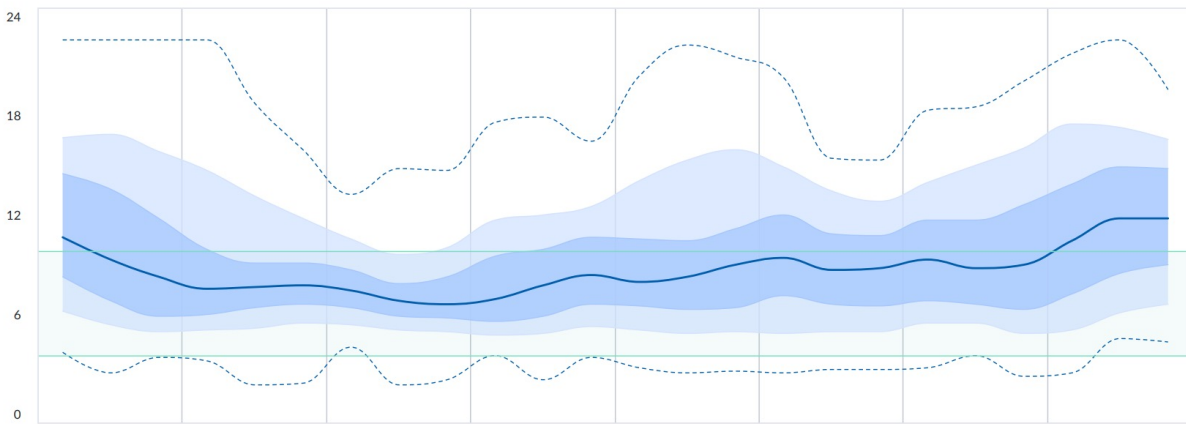


Summary

GMI	SD	3.7 mmol/L
7.4% (56.9 mmol/mol)	CV	39.2%
Average	Median	8.6 mmol/L
9.4 mmol/L	Highest	22.3 mmol/L
% Time CGM Active	Lowest	2.2 mmol/L
93.9% (28.2 days)		

Ambulatory Glucose Profile (AGP)

Target Range (3.9 - 10 mmol/L) 25 - 75% Lowest - Highest
Median 10 - 90%



Challenges of HCL in Dialysis

- Advanced complications
- High likelihood of other advanced complications – gastroparesis, advanced eye disease
- Potentially mental health issues
- Increased duration of insulin action
- Hypos are more challenging to treat
- HCL algorithms are not developed with PwD on dialysis
- Sensor accuracy?
- Very little guidance out there for using them in patients receiving haemodialysis

Haemodialysis with visual impairment

77 y/o White Caucasian Male

Type 1 diabetes from age 17

PMH of End-Stage Renal Failure

SPK in 2015 – both grafts failed in 2021

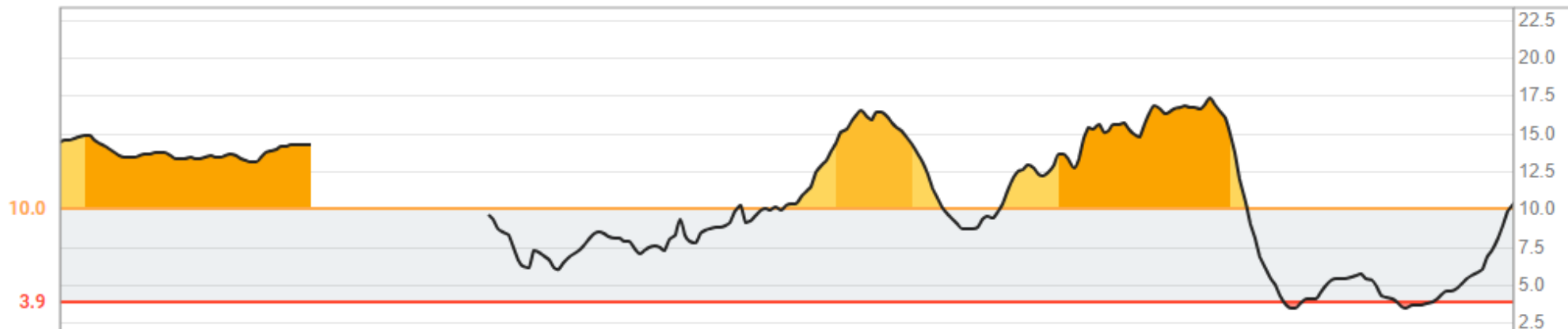
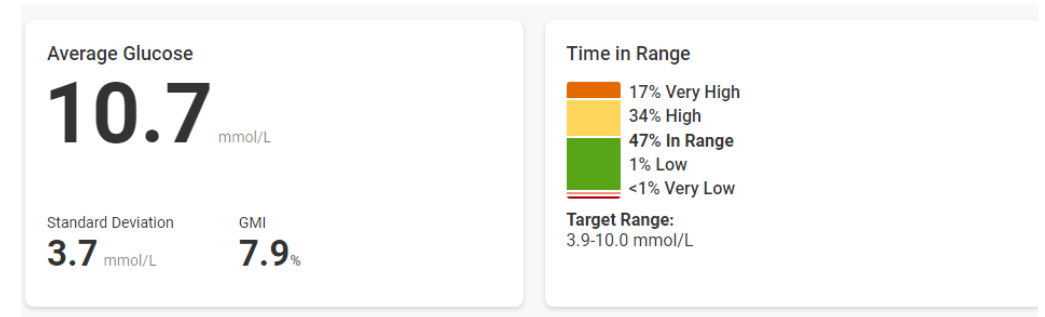
- Haemodialysis: M / W / F

MDI and CGM, Dexcom G6

Impaired hypo-awareness and severe hypoglycaemia history

Impaired vision and dexterity issues

Previously trialled Tandem T:slim X2 Insulin pump X

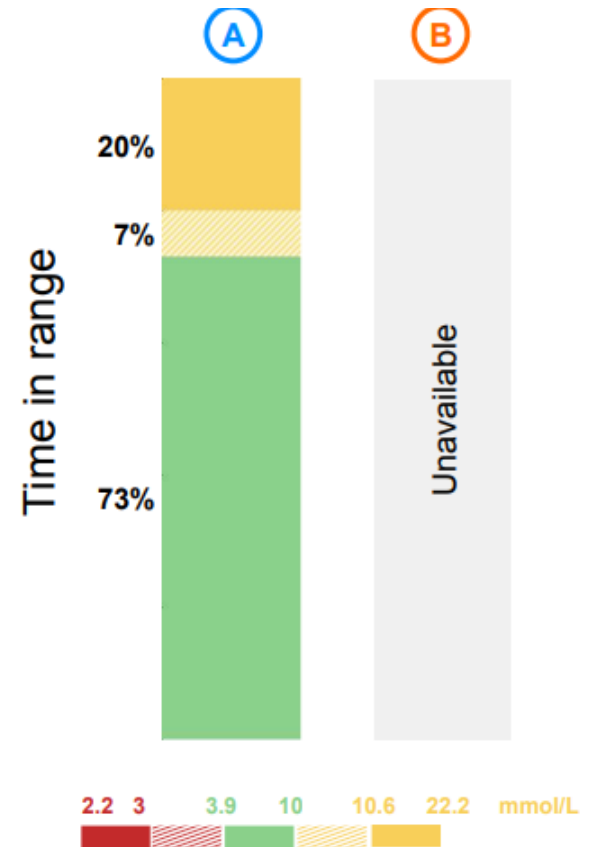


Post-HCL - Medtronic 780G

Improved time in range (73%)

Reduced hypoglycaemia (0%)

Reduced glucose variability



Thursday 22/02

TDD 26.0U

Total Basal 41% | 10.7U

Total Bolus 59% | 15.3U

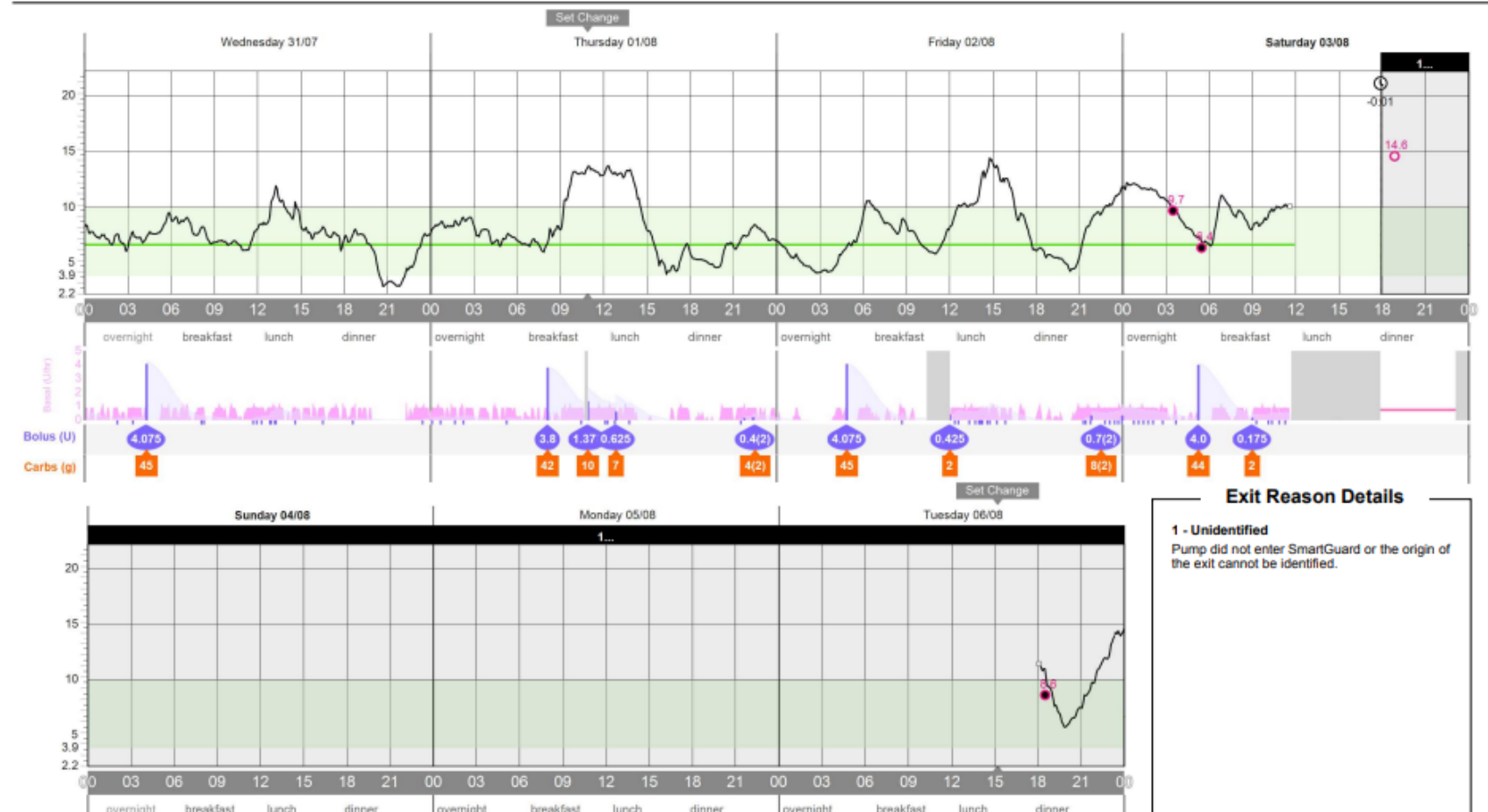
{ Bolus 86% | 13.1U

+ Auto Correction 14% | 2.2U }

Time in Range



- Sensor issues
- Post haemo-dialysis lows
- Challenges with adaptive basal when insulin sensitivity alters
- Active insulin time (vs duration of insulin action*)



*Walsh, Heinemann. JDST 2014 DOI: 10.1177/193229681351431

HDx - Pre HCL data

49 years old

Background

Type 1 diabetes, diagnosed in 2007.

On haemodialysis for end-stage renal disease.

Co-morbidities:

- Nonalcoholic fatty liver disease
- Chronic diarrhoea
- Hyponatraemia
- Lymphoedema
- Gastroparesis
- Mixed anxiety and depressive disorder
- Low back pain

Diabetic Complications:

- Proteinuric nephropathy
- Quiescent proliferative retinopathy
- Macular oedema
- Diabetic peripheral neuropathy
- Heel ulcer
- Diabetic foot ulcer

51 mmol/mol (March 2025)

Started on Medtronic insulin pump on 27/05/2025.

Medication

Alfacalcidol

Amlodipine

Atorvastatin

Creon (Pancreatin)

Descovy (Emtricitabine/Tenofovir)

Duloxetine

Eplerenone

Furosemide

GlucoRx

Metoclopramide

Ramipril

Sumatriptan

GLUCOSE STATISTICS AND TARGETS

20 February 2025 - 5 March 2025 14 Days

Time Sensor Active: 93%

Ranges And Targets For	Type 1 or Type 2 Diabetes
Glucose Ranges	Targets % of Readings (Time/Day)
Target Range 3.9-10.0 mmol/L	Greater than 70% (16h 48min)
Below 3.9 mmol/L	Less than 4% (58min)
Below 3.0 mmol/L	Less than 1% (14min)
Above 10.0 mmol/L	Less than 25% (6h)
Above 13.9 mmol/L	Less than 5% (1h 12min)
Each 5% increase in time in range (3.9-10.0 mmol/L) is clinically beneficial.	

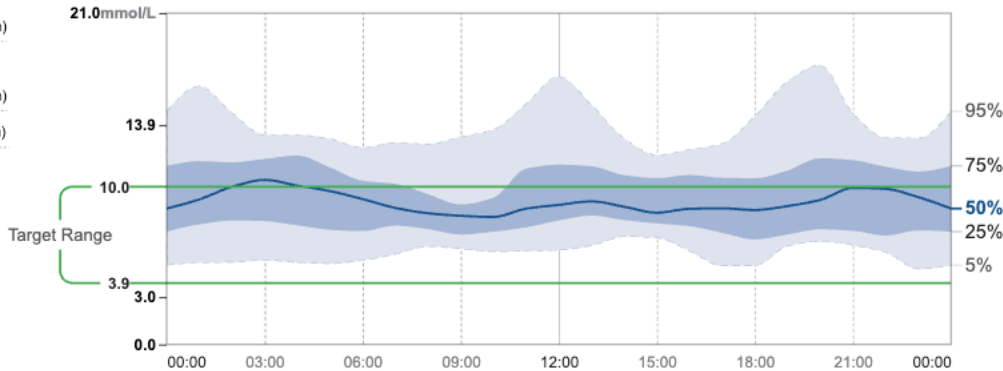
Average Glucose 9.3 mmol/L

Glucose Management Indicator (GMI) 7.3% or 56 mmol/mol

Glucose Variability 27.9%

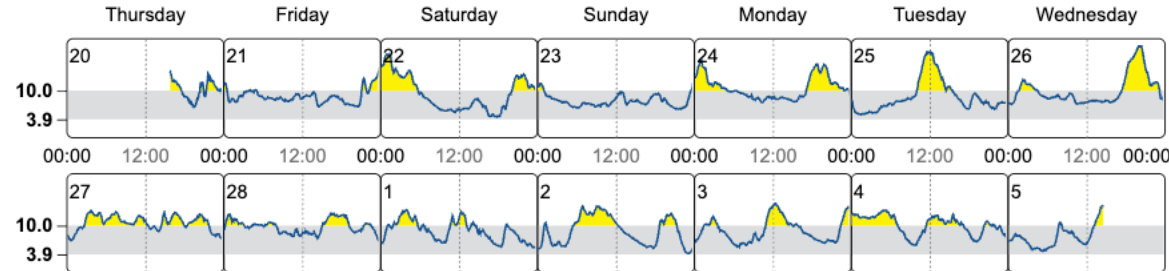
Defined as percent coefficient of variation (%CV); target ≤36%

TIME IN RANGES



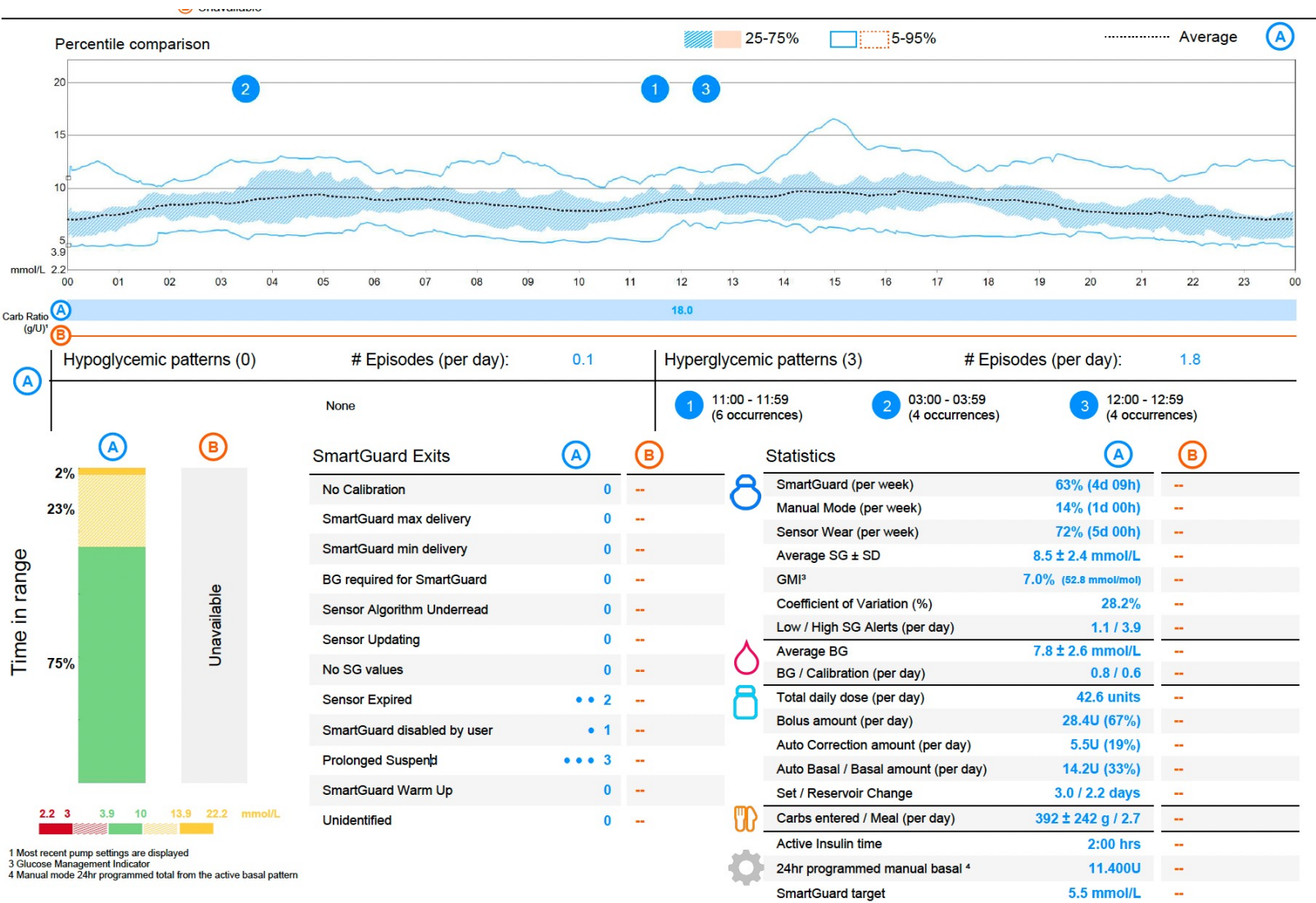
DAILY GLUCOSE PROFILES

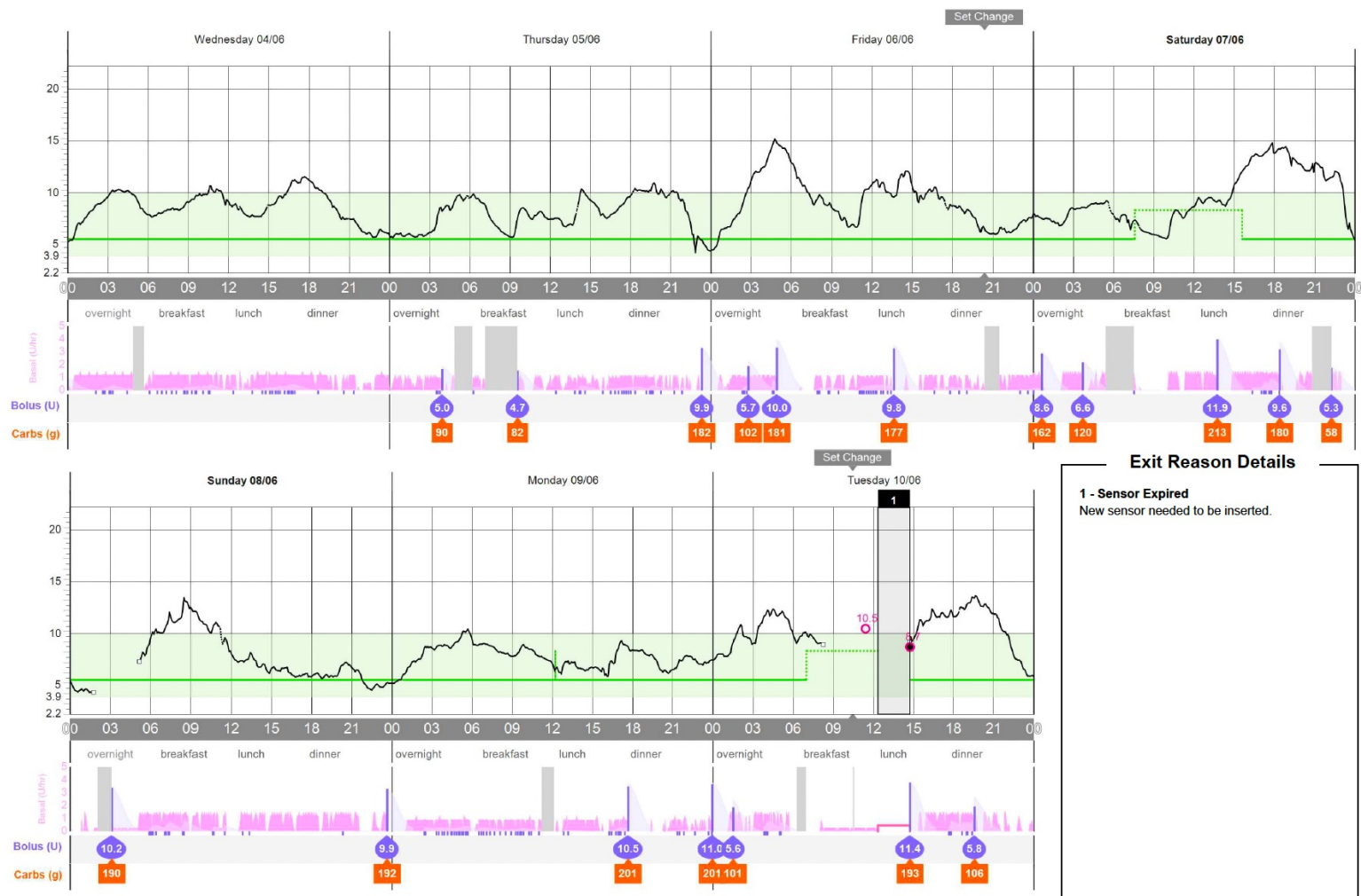
Each daily profile represents a midnight to midnight period with the date displayed in the upper left corner.



Source: Battelino, Tadej, et al. "Clinical Targets for Continuous Glucose Monitoring Data Interpretation: Recommendations From the International Consensus on Time in Range." Diabetes Care, American Diabetes Association, 7 June 2019, <https://doi.org/10.2337/dci19-0028>.

780G





AIT 2 hours
Glu target 5.5

Pre-HCL

48 yo Male originally from South Africa

Background

Type 1 diabetes since 2007

End-stage renal disease secondary to type 1 diabetes, now on haemodialysis.

HIV.

Gastroparesis.

Haemodialysis is undertaken on Mondays, Wednesdays, and Fridays

Past Medical History

Admission for peritonitis, which prompted a switch from peritoneal dialysis to haemodialysis in January 2024.

History of bilateral pneumonia, syphilis, and pleural effusion.

Diabetes History & Management

Pre-transplant insulin regimen included Levemir 22 units twice daily and NovoRapid, with Dexcom G7 monitoring.

Utilised Dexcom from 2024 to August 2024 prior to commencing hybrid closed-loop therapy.

Reviewed Omnipod 5 data from on and before 17/01/2025.

HbA1c (mmol/mol)

August 2024: 98

Medications

Alfacalcidol

Amlodipine

Aprepitant

Atorvastatin

Dolutegravir/TDF 50 mg

Dorzolamide

Doxazosin

Folic acid

Furosemide

Irbesartan

Lamivudine

Ondansetron

Quinine

Salbutamol

Senna

Sertraline

Sevelamer

Sildenafil

Dialysis-Related Medications

Mircera injections

Dalteparin

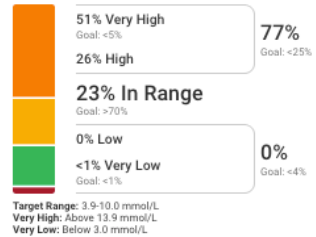
Pre-HCL

Dexcom G6 + MDI

dexcom | capturAGP[®] v5.0

Time in Ranges Goals for Type 1 and Type 2 Diabetes

Each 5% increase in the Target Range is clinically beneficial.
Each 1% time in range = about 15 minutes per day



Emile Cupido

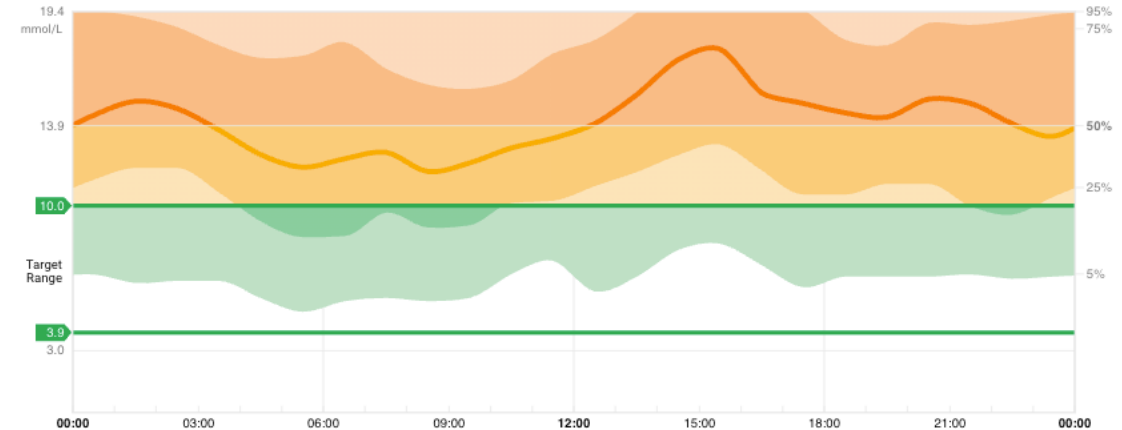
DOB: 3 March 1977

Glucose Metrics

Average Glucose Goal: <8.5 mmol/L	14.4 mmol/L
GMI Goal: <7%	9.5%
Coefficient of Variation Goal: <36%	35.8%
Time CGM Active	99.1%

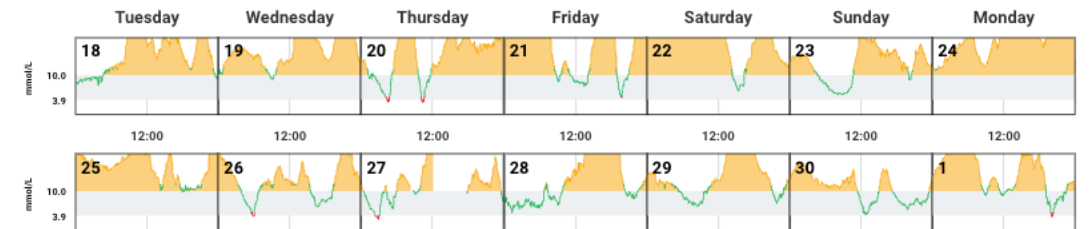
Ambulatory Glucose Profile (AGP)

AGP is a summary of glucose values from the report period, with median (50%) and other percentiles shown as if they occurred in a single day.



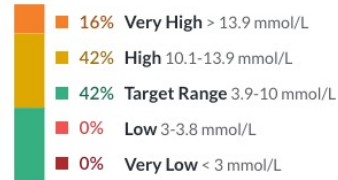
Daily Glucose Profile

Each daily profile represents a midnight-to-midnight period.



Started on OP5

Glucose (CGM)



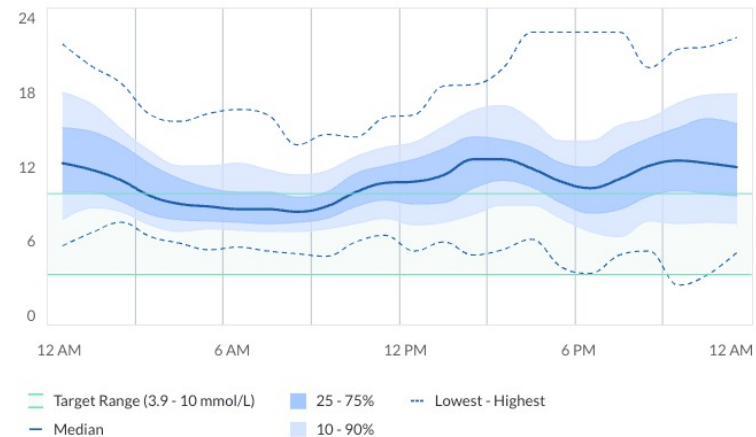
% Time CGM Active 94.2% (28.3 days)

GMI ?	8.1% (64.6 mmol/mol)
Average	11 mmol/L
SD	3 mmol/L
CV	27.6%
Median	10.6 mmol/L
Highest	HI mmol/L
Lowest	3 mmol/L

AGP

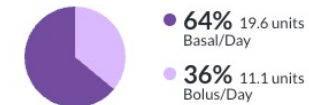
Glucose (mmol/L)

[What is AGP?](#)



Insulin - Device ?

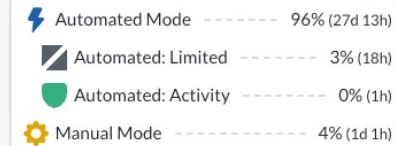
From Insulin Pump



Insulin/Day	30.7 units
Overrides (%)	0% (0 boluses)
# Bolus/Day	3.7

System Details

Insulet Omnipod® 5 System (28d 15h)



Diet

Dec 19, 2024 - Jan 17, 2025

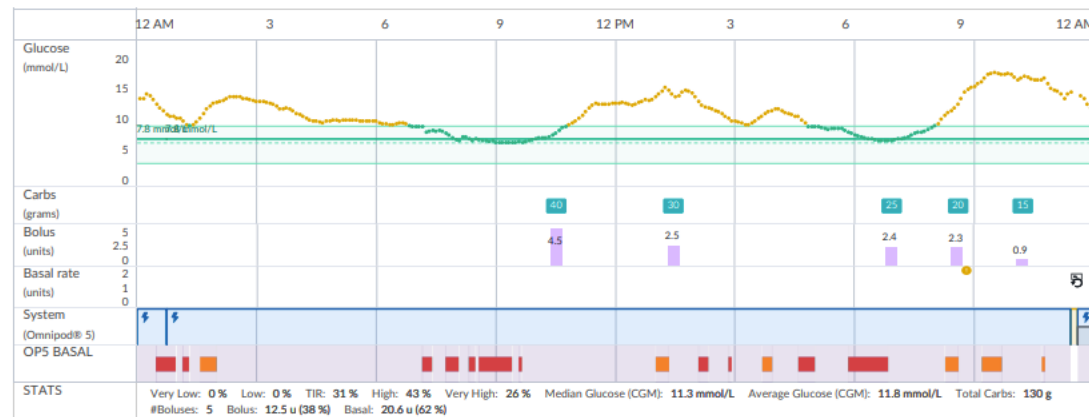
115.7 g Carbs/Day 3.9 Entries/Day

Activity

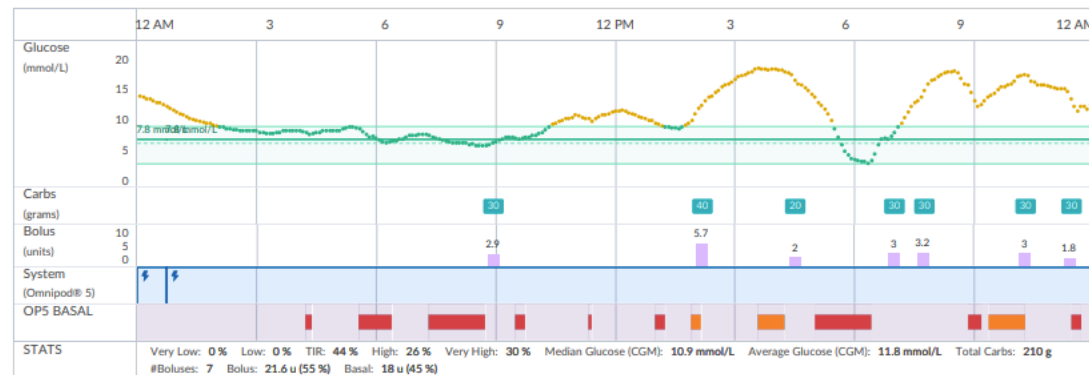
Dec 19, 2024 - Jan 17, 2025

OP5 with HDx MWF

Tuesday, January 14, 2025



Monday, January 13, 2025



Sunday, January 12, 2025

Insulet Omnipod® 5

System

Controller Device: -
Serial Number: 06040000-000051368

Device 1 of 1

General

Active Insulin Time: 4 hours

Basal

Max Basal Rate: 1.7 U/hour
Temporary Basal Enabled: ON
Active basal program: Basal 1

Bolus

Min BG for Bolus Calc: 3.9 mmol/L
Extended Bolus: ON
Reverse Correction: ON
Max Bolus: 8 U

Basal

Basal 1 Active
12:00 AM (24 hr): 0.7 Units/hr
Total: 16.8 Units

Insulin : Carb Ratios

Profile Active
12:00 AM (24 hr): 10 g/Unit

Sensitivity (ISF, Correction)

Profile Active
12:00 AM (24 hr): 3.2 mmol/L

BG Target Range

Profile Active
12:00 AM (24 hr): 7.8 mmol/L (+0/-0)

BG Correction Threshold

Profile Active
12:00 AM (24 hr): 7.8 mmol/L

Current status

HbA1c (mmol/mol)

August 2024: 98

October 2024: 51

January 2025: 57

NB- urgent eye screening!

Post-simultaneous pancreas-kidney (SPK) transplant 8 months ago

Off all insulin therapy following successful SPK transplant.

HbA1c (mmol/mol)

July 2025: 32

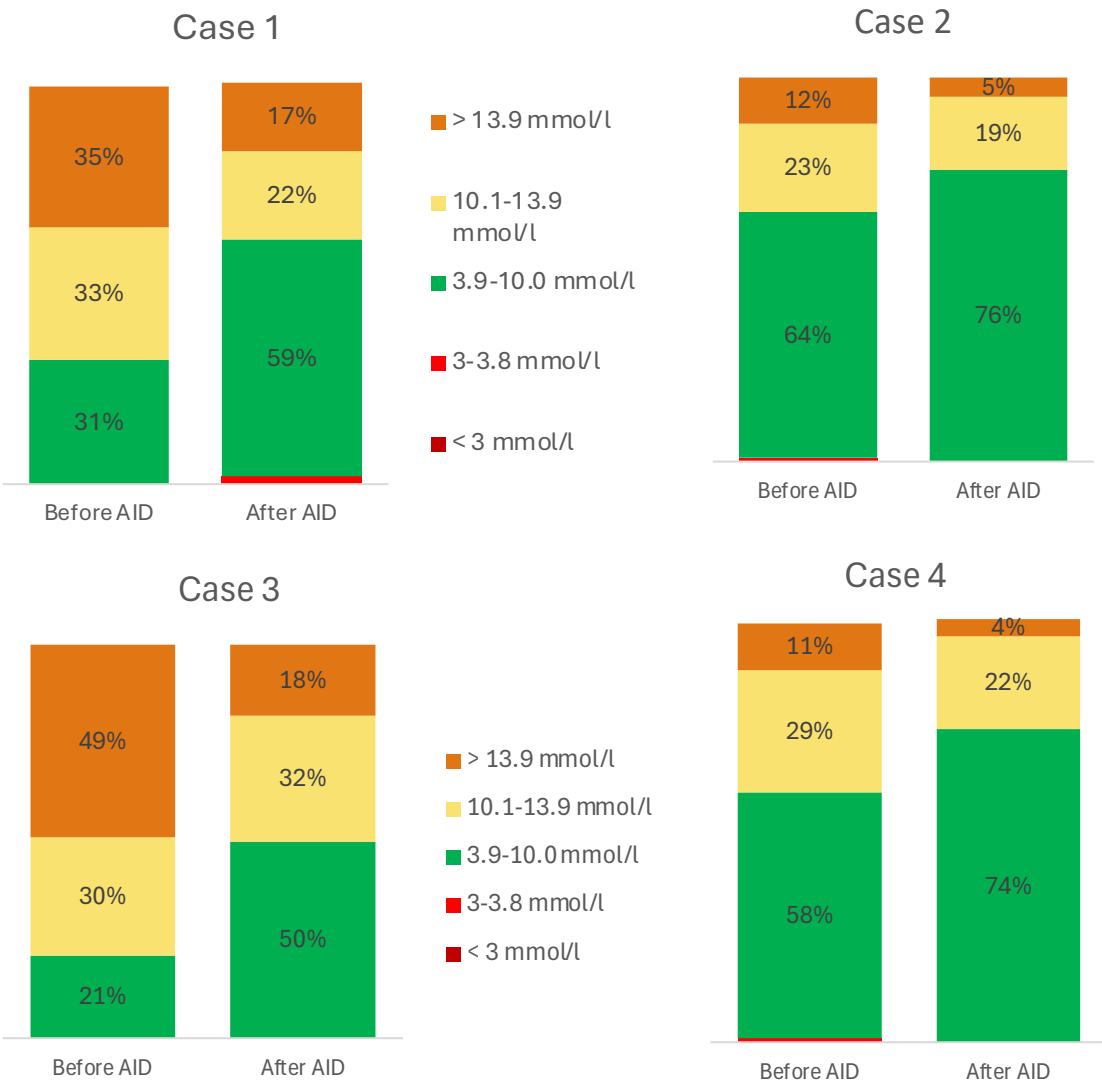
September 2025: 33

Haemodialysis and HCL

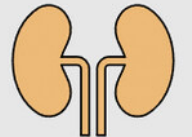
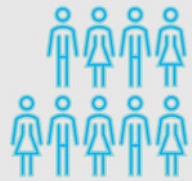





Case series of 4 adults with T1D on haemodialysis

- 3 on Medtronic 780G
- 1 on CamAPSFX
- Improvement in glycaemic metrics and positive feedback

Chaudhry, K., Hyslop, R., Johnston, T., Pender, S., Hussain, S., & Karalliedde, J. (2024). *Diabetes Research and Clinical Practice*, 111800



Using automated insulin delivery to address the clinical challenges of glycaemic management in people with type 1 diabetes and kidney failure on maintenance haemodialysis

Study Population	Methods	Key Results
<p>We studied 9 individuals (5 women, 4 men) with T1D and ESKD, all on a stable thrice-weekly haemodialysis regimen. Participants were recruited from specialist multidisciplinary diabetes clinics at three university hospitals in London.</p> <div></div> <p>The mean age was 39.9 years (range 33–50), mean T1D duration of 30.7 years (range 17–48). All had prior experience with diabetes technology (CGM and/or insulin pump therapy). Median follow up was 7 months.</p>	<ul style="list-style-type: none">Initiated one of several commercially available AID systems: Medtronic 780G (with Guardian 4 sensor), Tandem T-Slim (with Dexcom G6), and Omnipod 5 (OP5) (with Dexcom G6 or Freestyle Libre 2+)Pre-AID and post-AID data were obtained from CGM, electronic health records (EHR) and/or pump downloadsThe primary outcome was change in time in range (TIR) (3.9-10 mmol/L) following AID initiation <div></div>	<ul style="list-style-type: none">Improvement in mean time in TIR with AID from 39.7 (±17.1) % to 59.8 (±18.1) % (p = 0.001)Mean glucose reduced from 13.1 (±2.4) mmol/L to 9.8 (±1.9) mmol/L (p = 0.0002).Glucose variability improved from 39.8 (±3.8) % to 33.8 (±6.5) % (p = 0.01).Dialysis-day TIR increased from 32.7 (±17.7) % to 62.3 (±14.9) % (p = 0.002).HbA1c improved from 78.6 (±12.8) mmol/mol to 56.1 (±8.5) mmol/mol (p = 0.003)Mean time below range (TBR) (< 3.9 mmol/L) fell from 4.0 (±5.4) % to 1.4 (±1.3) % (P = 0.1)AID was safe and well tolerated <div><div><div>TIR (+20.1%)</div></div><div><div>TIR on dialysis days (+59.6%)</div></div><div><div>Mean glucose (-3.3 mmol/L)</div></div></div>

Peritoneal dialysis?

- **44yo with 23 y T1D**
- **Past Medical History**
 - Type 1 diabetes mellitus with multiple complications:
 - Diabetic macular oedema
 - End-stage renal failure (on peritoneal dialysis)
 - Autonomic neuropathy
 - Gastroparesis
 - Quiescent proliferative retinopathy (previous pan-retinal photocoagulation and vitrectomy)
 - Peripheral nerve disease
 - Essential hypertension
 - Gastro-oesophageal reflux
 - Hypercholesterolaemia
 - Obesity
 - Recurrent painful ophthalmoplegic neuropathy
 - Posterior vitreous detachment
 - History of transient cerebral ischaemia (2019)

Current Medications

- Doxazosin 2 mg
- Loperamide 2 mg
- Metoclopramide
- Bisoprolol
- Losartan
- Atorvastatin
- Indapamide
- NovoRapid insulin
- Sildenafil
- Alprostadil
- Hydrocortisone
- Clopidogrel
- Renavit one tablet daily

Recent History & Investigations

- Ophthalmology:

- Mild recurrence of macular oedema in 2023 and 25

- Renal:

- PDx April 2025

- SPK list

- switch to daytime dialysis (four times a day)

- Diabetes/General:

- HbA1c trend:

- 25/03/2025: 55 mmol/mol

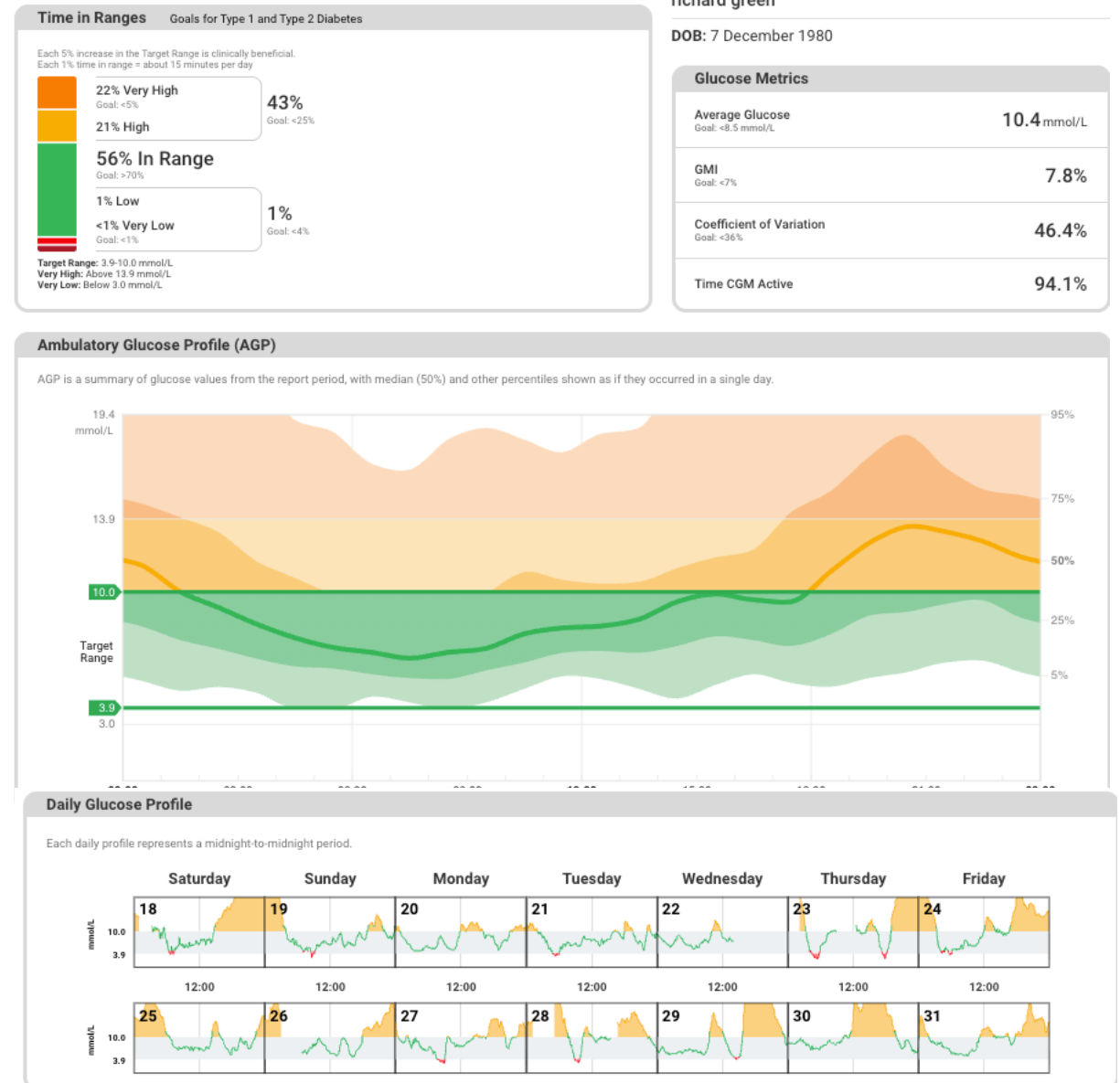
- 17/12/2024: 57 mmol/mol

- 18/06/2024: 60 mmol/mol

- 16/02/2024: 60 mmol/mol

- On MDI and Dexcom

- Started tandem insulin pump 26/04/2025.



HCL

Glucose (CGM)



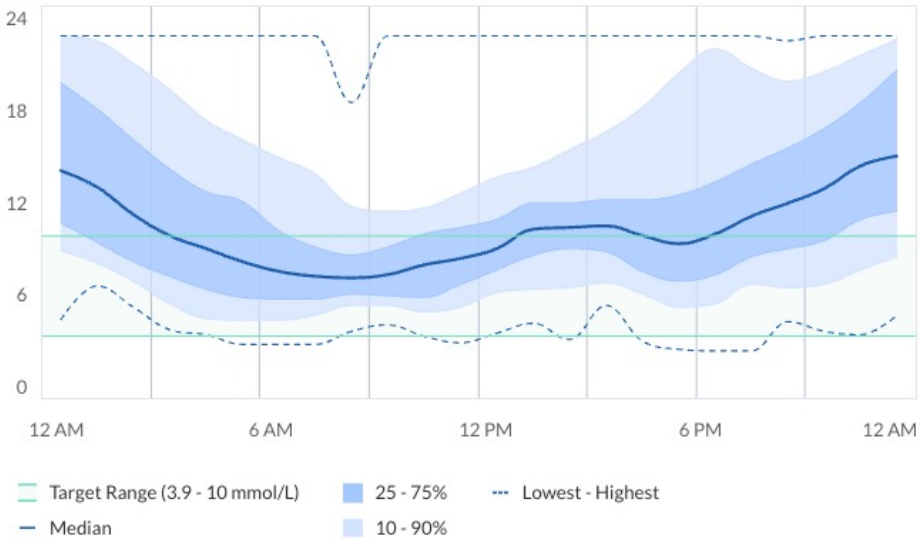
% Time CGM Active 96.5% (29 days)

GMI ?	8.1% (64.8 mmol/mol)
Average	11.1 mmol/L
SD	4.7 mmol/L
CV	42.9%
Median	9.9 mmol/L
Highest	HI mmol/L
Lowest	2.9 mmol/L

AGP

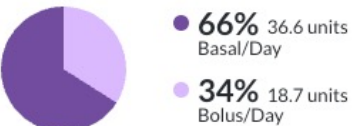
Glucose (mmol/L)

[What is AGP?](#)



Insulin - Device ?

From Insulin Pump



Insulin/Day	55.3 units
Automatic Bolus/Day	14.8 units (27%)
Overrides (%)	14% (52 boluses)
# Bolus/Day	12.8
# Automatic Bolus/Day	11.1

System Details

Tandem t:slim X2 (29d 2h)



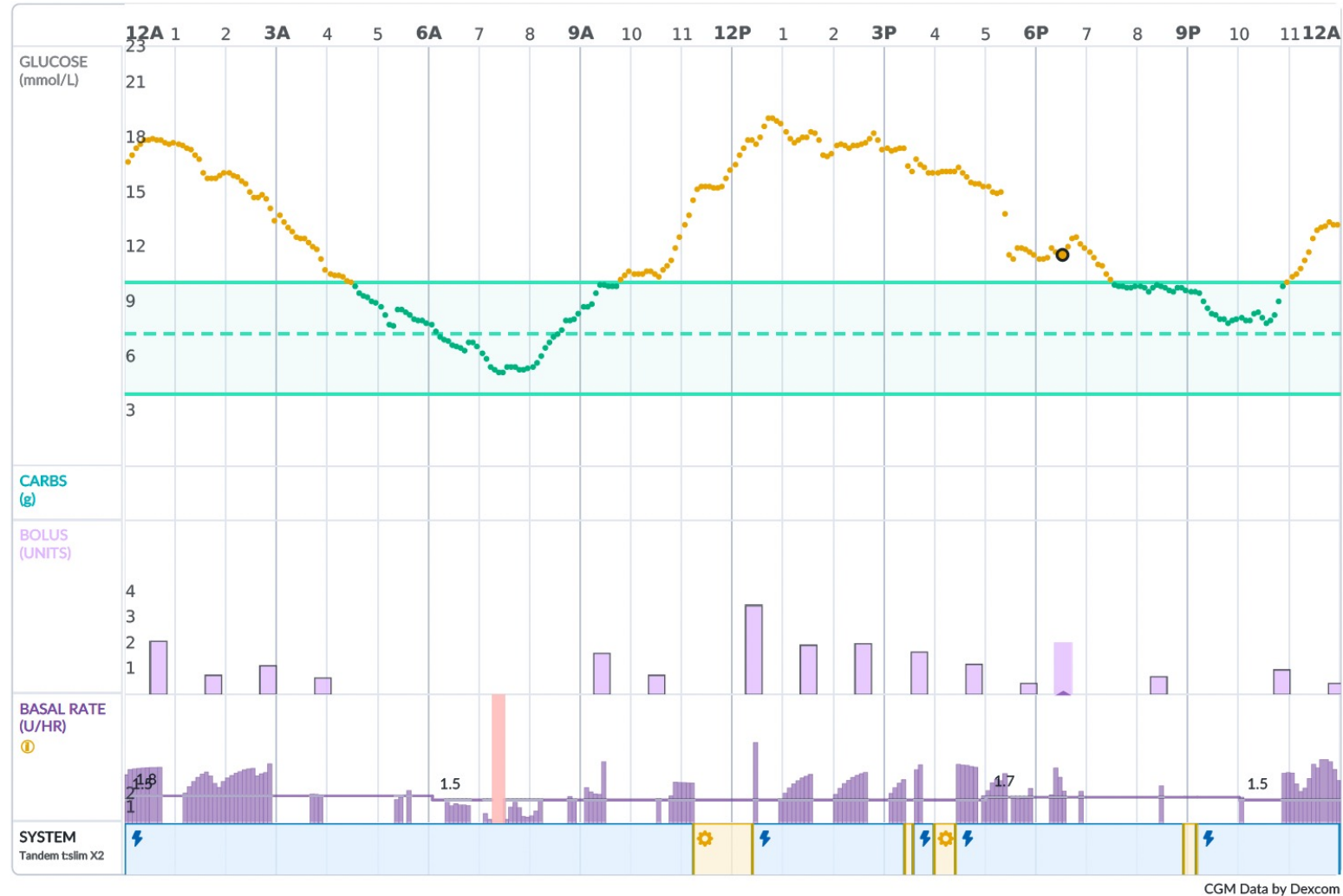
Control-IQ	92% (26d 20h)
Activity - Sleep	0%
Activity - Exercise	0%
Manual	8% (2d 6h)

LGS/PLGS

Time Suspended/Day	53m
Avg Suspensions/Day	2.4
Avg Suspension/Time of Day	Morning - 36% Afternoon - 11% Evening - 19% Night - 34%

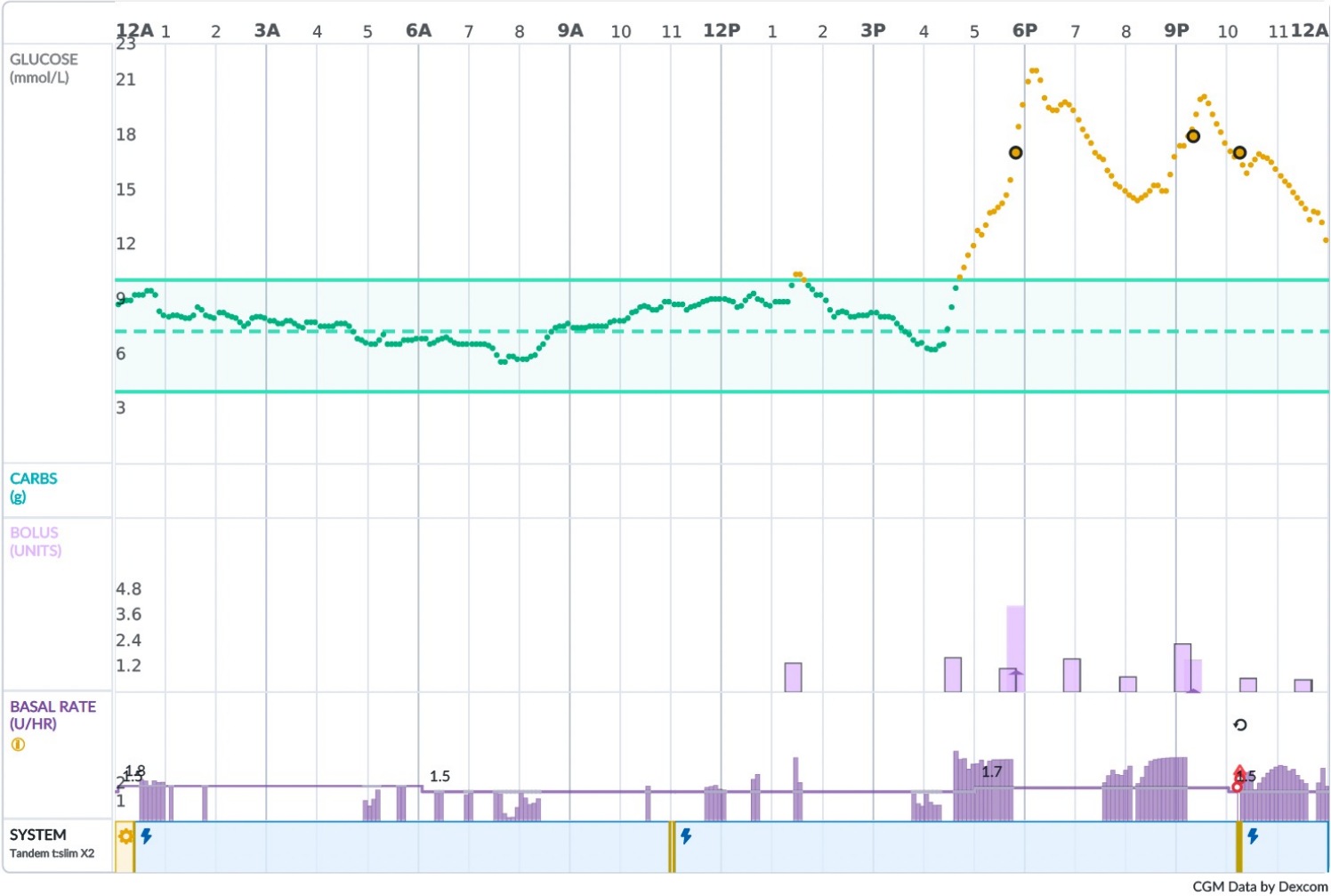
[Show more](#)

Difficult!



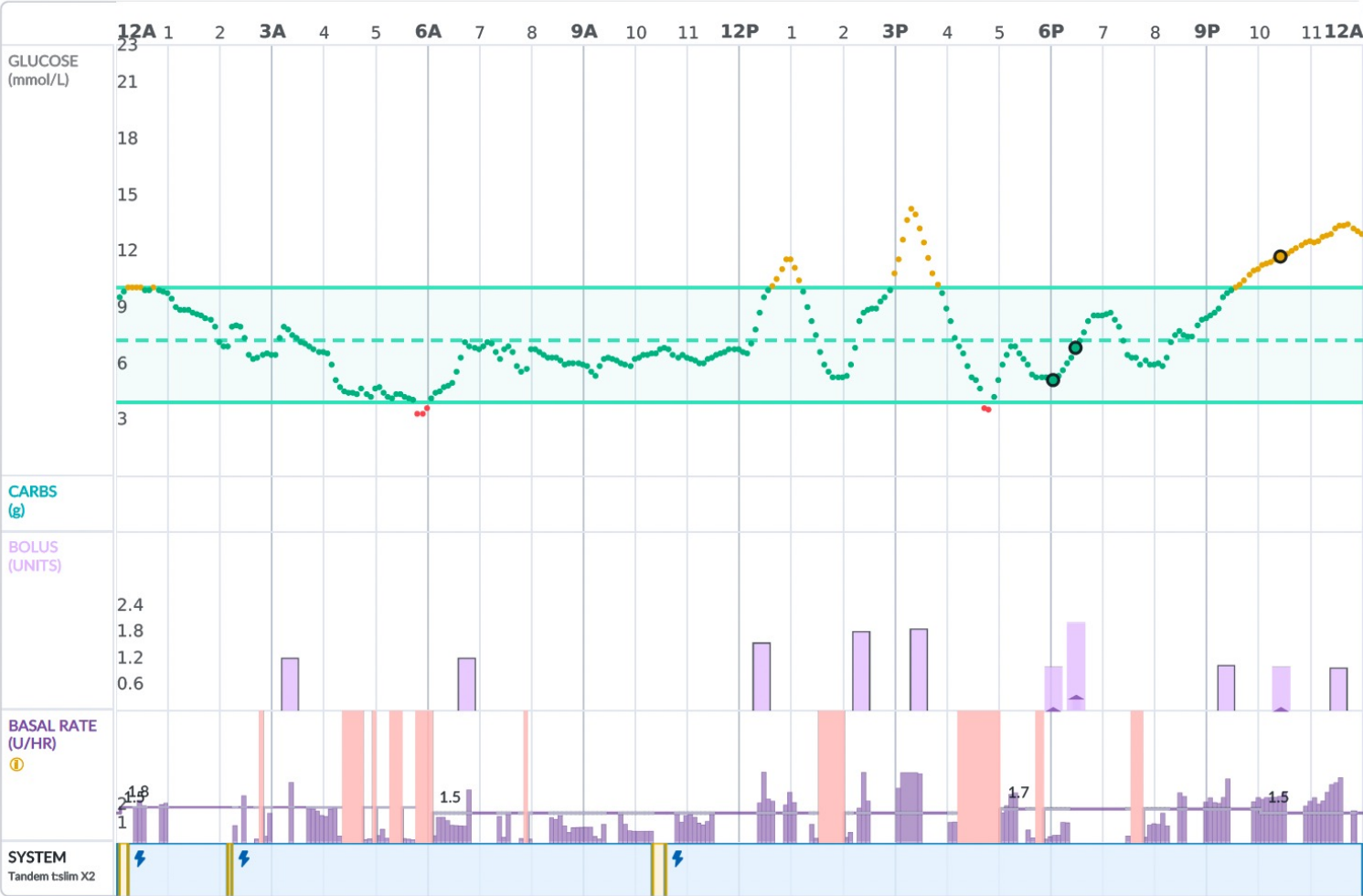
Still difficult!

Tuesday, Sep 23, 2025



Progress?

Sunday, Oct 12, 2025



HbA1c 14/10/2025: 53 mmol/mol

Profile 1 Active	Basal Units/hr	Sensitivity (ISF, Correction) mmol/L	Insulin : Carb Ratios g/Unit	BG Target Range mmol/L
12:00 AM (6 hr)	1.8	2	10	6.1 (+0/-0)
6:00 AM (5 hr)	1.5	2	10	6.1 (+0/-0)
11:00 AM (6 hr)	1.5	2	10	6.1 (+0/-0)
5:00 PM (5 hr)	1.7	2.5	10	6.1 (+0/-0)
10:00 PM (2 hr)	1.5	2.5	10	6.1 (+0/-0)
Total	38.8 Units			

Pump Boluses

Bolus Table only shows the first 10 bolus events for the day.

Statistics

Learning points: HCL in HDx

HCL is beneficial at increasing TIR in this high risk group of people
However it is not without challenges

Nil adverse events in a high risk population e.g. DKA or severe hypoglycaemia

Freedom from multiple insulin injections

Tips to consider

Conservative treatment targets / AIT settings
Avoid bolus dosing during dialysis sessions

780G

- AIT 4 hours and Glucose target 6.7
- Activity mode may not be needed beyond first few weeks
- Remote connectivity key

CamAPS FX

- Glucose target 7-7.5
- Avoid boost after HDx
- Ease off if concerns first few HDx sessions but avoid (as will not adapt in future)

OP5

Glucose target 7.8-8.3

Control IQ

More experience needed but potential to influence ISF / basal
Profile changes between HDx/Non HDX can be challenging for user group, activity mode consideration if initial concerns
Remote Tandem Diabetes Source needed

Further tips

- MDT teamwork key - with regular input upon initiation
- Capacity and training issues
 - May need 1:1 start
 - Daily contact
 - Review supervised set change
 - Bring back in clinic at 4 wks for review
- Eye assessment
- Close liaison with Renal satellite unit

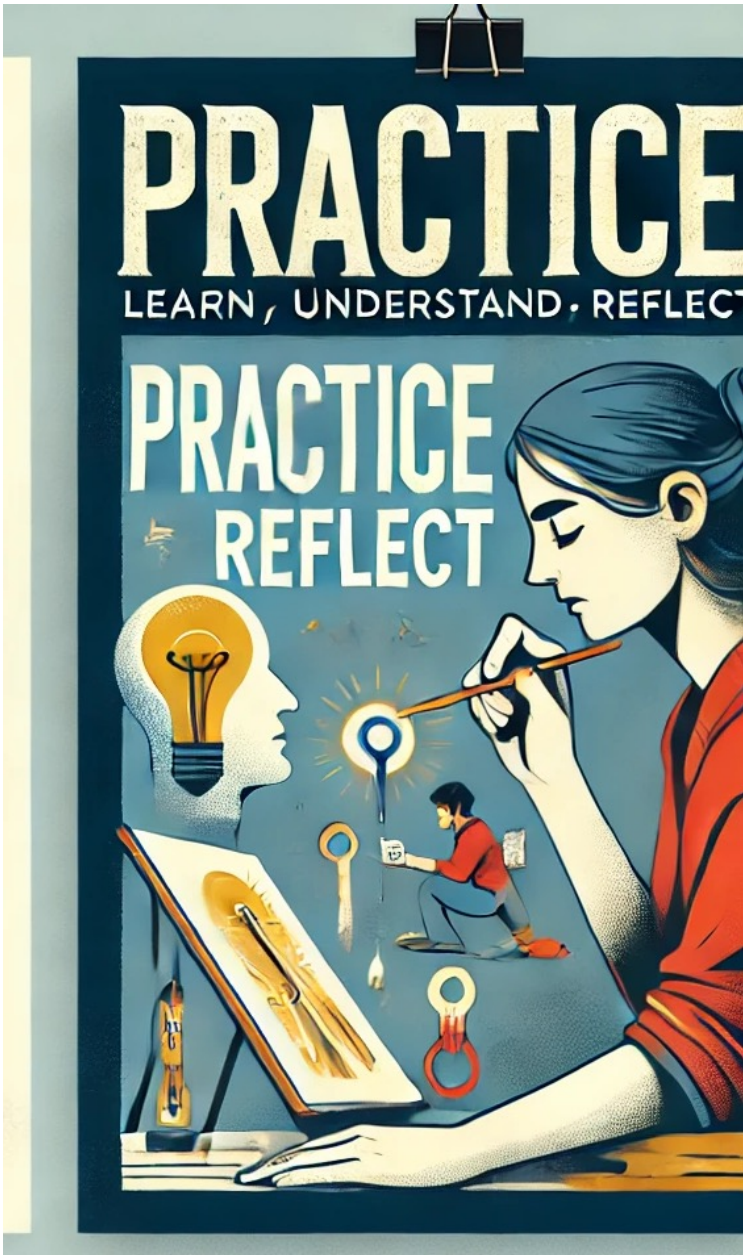
Practical considerations for AID in haemodialysis

- **Assess suitability:**
 - Before starting dialysis, individuals using AID systems should be assessed to confirm whether AID therapy can be continued safely and to determine if any setting adjustments are needed.
- **Coordinated Care Plan:**
 - If the patient is considered appropriate to remain on AID therapy, a multidisciplinary diabetes care plan should be created and recorded within the shared electronic medical record.
 - When beginning AID treatment, structured follow-up is essential, typically involving 2–3 contacts per week (face-to-face or virtual) for about the first two weeks. After this initiation phase, appointments can be reduced to weekly as needed. Once therapy has been established for two months, the review schedule may be tailored to individual requirements
 - Consider the presence of diabetes complications such as advanced retinopathy and neuropathy (peripheral and autonomic) which can be exacerbated by rapid fluxes in glycaemic control. Prior to AID initiation ensure the patient is up to date with retinopathy screening and there is no active eye disease which requires treatment. In the context of such complications, initiation of AID may be postponed, or conservative glycaemic treatment goals may need to be adopted. Using an insulin pump in manual mode for the initial period may be a viable strategy, with activation of AID at a later stage to ensure gradual and cautious glycaemic improvement.
- **Recommendations for AID Use During Dialysis:**
 - Modify blood glucose target, active insulin time and basal profile to more conservative settings when starting dialysis or initiating AID in the context of end-stage kidney disease. SmartGuard™ and SmartAdjust™ provide glucose target options as high as 8.3 mmol/L, Control IQ 8.9 mmol/L and CamAPS™ even higher (up to 11 mmol/L). Glucose target of 8.3 mmol/l at initiation would be suitable in most cases in the context of ESKD.
 - Advise patients to refrain from administering bolus doses during dialysis to due to risk of hypoglycemia.
 - Avoid paracetamol when using Medtronic CGM sensors, as it can interfere with sensor accuracy.
 - Keep the control device (e.g., smartphone) within six meters of the person at all times.
 - If low blood glucose is a concern during dialysis sessions, consider activating temporary modes (activity/exercise/sleep) to increase target glucose levels for a designated time period. Activation 1-2 hours prior to dialysis continuing for 1-2 hours after dialysis (or longer depending on individual susceptibility to hypoglycaemia) could be a viable strategy.

The background image is a collage of financial and data-related elements. It includes a line graph with a grid, a calendar showing days of the week (Mon, Tue, Wed, Thu, Fri, Sat), a calculator with numbers, and various other data visualizations like bar charts and pie charts. The overall color scheme is dark with blue and orange accents.

What more is needed? Research studies

- Studies in populations with complications - to aid optimal device design, algorithm development, features and simplicity
- Trials to help demonstrate safety and efficacy for licensing and remuneration
- Guidance and consensus from HCPs
- **Pseudocode from industry to guide use in complex scenarios**



Summary

HCL can offer benefits but need understanding, **teamwork**, adaptations and tailored advice in complex settings

Learn by experience

Discuss cases as teams

Manage as multidisciplinary teams

Share learnings in regional networks



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Pioneering better health for all

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