

LA TECNOLOGIA PER LA CURA DEL DIABETE

Overview della tecnologia nel diabete

Dott. Federico Boscari

UOC Malattie Del Metabolismo

Azienda Ospedale Università di Padova

DALLA SMART MDI

ALL'HYBRID CLOSED LOOP

ROMA 1-2 Aprile 2025

Il dr. **Federico Boscari** dichiara di aver ricevuto negli ultimi due anni compensi o finanziamenti dalle seguenti Aziende Farmaceutiche e/o Diagnostiche:

- Movia
- Roche Diabetes Care
- Sanofi Aventis
- Glooko
- Tandem
- Novo Nordisk

Dichiara altresì il proprio impegno ad astenersi, nell'ambito dell'evento, dal nominare, in qualsivoglia modo o forma, aziende farmaceutiche e/o denominazione commerciale e di non fare pubblicità di qualsiasi tipo relativamente a specifici prodotti di interesse sanitario (farmaci, strumenti, dispositivi medico-chirurgici, ecc.).

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16.30 Overview della tecnologia nel diabete

Federico Boscari

16.50 Selezione del paziente e comunicazione efficace: i vantaggi dell'Hybrid Closed Loop verso la terapia MDI

Elena Succurro

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ALL'HYBRID CLOSED LOOP

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Nuzzo, M.G.; Schettino, M. Advanced Technology (Continuous Glucose Monitoring and Advanced Hybrid Closed-Loop Systems) in Diabetes from the Perspective of Gender Differences. *Diabetologia* 2023, 4, 519-526. <https://doi.org/10.3390/diabetology4040045>



diabetes and technology



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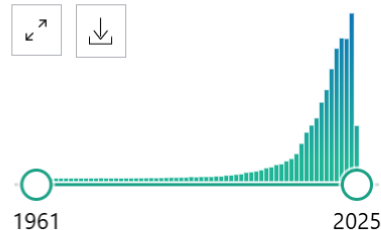
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72,330 results

« < Page 1 of 7,233 > »

RESULTS BY YEAR



PUBLICATION DATE

- ☐ 1 year
- ☐ 5 years
- ☐ 10 years

☐ **Digital health technology and diabetes management.**

1 Cahn A, Akkew A, Rasi.

Cite J Diabetes. 2018 Jan;10(1):10-17. doi: 10.1111/1753-0407.12606. Epub 2017 Nov 6.

PMID: 28871765 Review.

Share **Diabetes** care is largely dependent on patient self-management and empowerment, given that patients with **diabetes** must make numerous daily decisions as to what to eat, when to exercise, and determine their insulin dose and timing if required. ...Better understanding ...

☐ **Diabetes Technology and the Human Factor.**

2 Liberman A, Barnard K.

Cite Diabetes Technol Ther. 2019 Feb;21(S1):S138-S147. doi: 10.1089/dia.2019.2511.

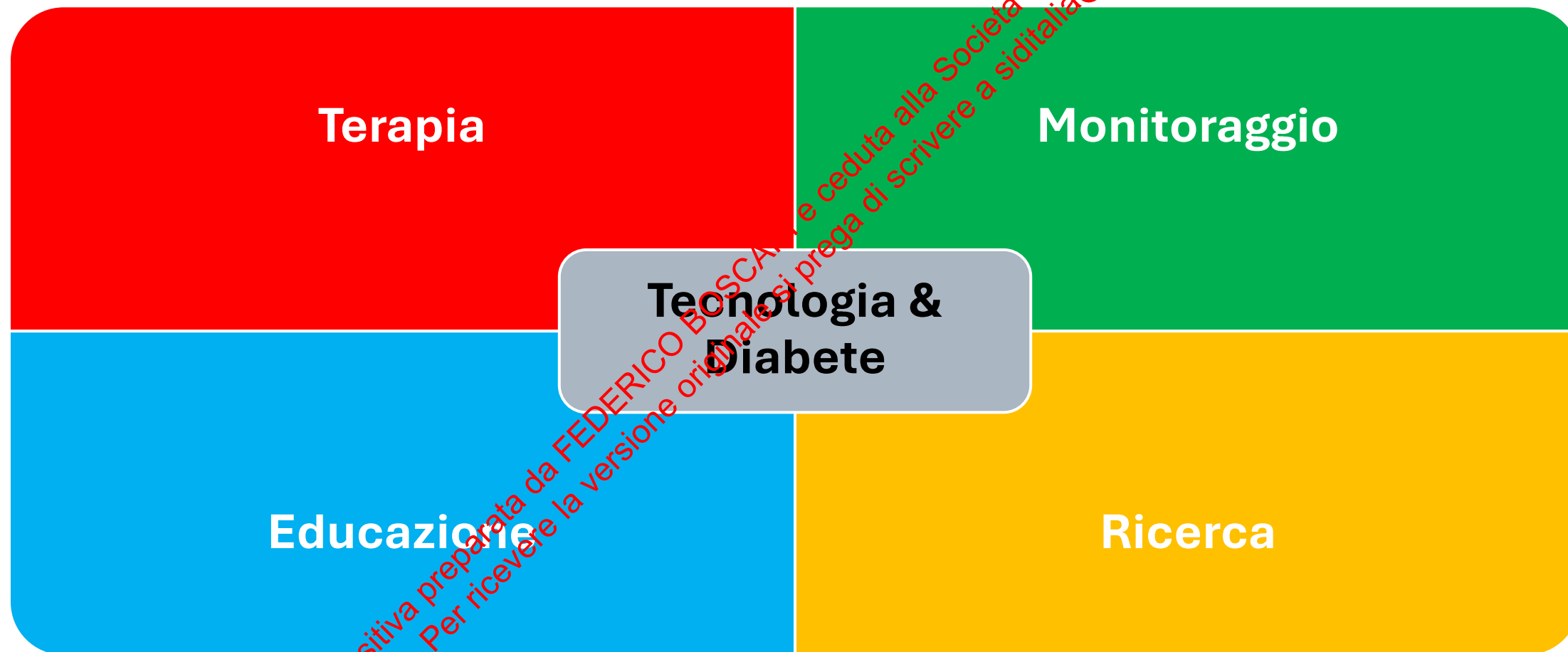
PMID: 30785329 Review. No abstract available.

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In che modo la tecnologia interviene nella gestione del diabete

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In che modo la tecnologia interviene nella gestione del diabete



In che modo la tecnologia interviene nella gestione del diabete

Terapia

**Tecnologia &
Diabete**

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Somministrazione di insulina

- Microinfusori di insulina
- Sistemi automatici per la somministrazione di insulina
- Smart Pen

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AID (Automated Insulin Delivery)

Struttura generale

Sistemi ibridi

Similitudini

Peculiarità

Tipo di sensore

Tipo di microinfusore

Algoritmo

Condivisione dati

Indicazioni cliniche

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AID

Outcome	No. of comparisons	No. of patients		Mean difference or relative risk (95% CI)	P value
		Intervention	Control		
Time in range	62	2,292	2,008	11.74 (9.37 to 14.12)	<0.001
Time below range	60	2,242	1,878	-1.20 (-1.62 to -0.79)	<0.001
Time above range	54	1,979	1,684	-10.17 (-13.18 to -7.16)	<0.001
Coefficient of variation, %	41	1,682	1,402	-1.31 (-2.16 to -0.47)	0.002
HbA1c, %	25	1,453	1,171	-0.36 (-0.44 to -0.28)	<0.001
Glycemia risk index	6	139	139	-3.74 (-6.34 to -1.14)	0.005
Insulin dose	44	1,508	1,376	0.03 (-0.08 to 0.15)	0.577
Severe hypoglycemia	21	1,134	977	0.92 (0.65 to 1.31)	0.661
Diabetic ketoacidosis	9	554	445	1.15 (0.47 to 2.80)	0.766

CI, confidence interval; HbA1c, glycosylated hemoglobin.

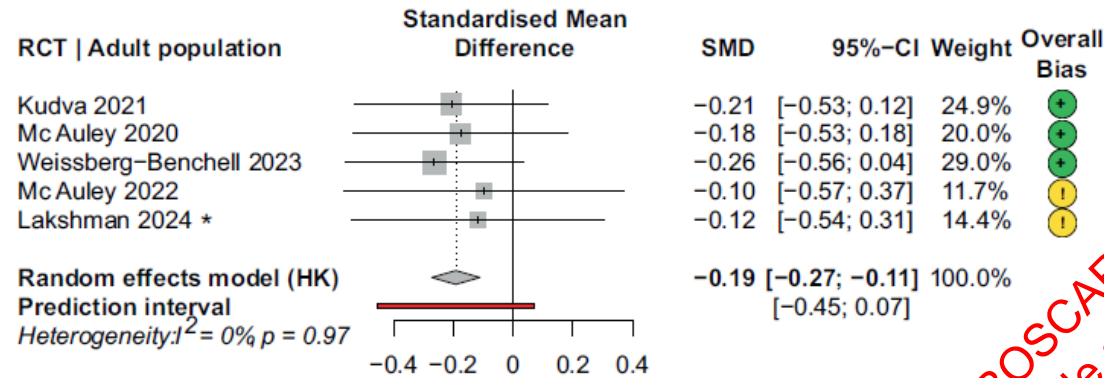
Fan W, Deng C, Xu R, Liu Z, Leslie RD, Zhou Z, Li X. Efficacy and Safety of Automated Insulin Delivery Systems in Patients with Type 1 Diabetes Mellitus: A Systematic Review and Meta-Analysis. Diabetes Metab J. 2025 Mar;49(2):235-251. doi: 10.4093/dmj.2024.0130. Epub 2024 Nov 13. PMID: 39533812.

Outcomes and subgroups	No. of comparisons	No. of patients		Mean difference between AID systems and other insulin therapy (95% CI), %	P value for overall effect	P value for subgroup differences	Weight, %	I ² , %
		Intervention	Control					
Time in range, %								
Total comparisons	62	2,292	2,008					
Timing of intervention								
24 hours	60	2,251	1,968	11.812 (9.396 to 14.229)	<0.001	0.479	97.22	94.8
Overnight	2	41	40	9.485 (3.516 to 15.454)	0.002		2.78	0.0
Hormone								
Single	54	2,107	1,822	11.021 (8.495 to 13.549)	<0.001	0.056	87.59	95.0
Dual	8	185	186	16.950 (12.412 to 22.488)	<0.001		12.41	79.6
Age								
Adults	23	814	713	15.107 (8.815 to 17.400)	<0.001	0.245	36.68	93.3
Children and adolescents	26	834	658	11.171 (8.191 to 15.551)	<0.001		41.68	95.5
Mixed population	13	644	527	8.933 (5.819 to 12.048)	<0.001		21.64	82.7
Baseline HbA1c, %								
<7.5	21	660	661	8.844 (5.906 to 11.782)	<0.001	0.039	34.13	91.6
≥7.5	41	1,632	1,347	13.245 (10.270 to 16.220)	<0.001		65.87	92.9
Diabetes duration, yr								
<20	38	1,543	1,258	11.217 (8.443 to 13.991)	<0.001	0.555	64.51	94.2
≥20	21	707	708	12.756 (8.470 to 17.041)	<0.001		35.49	93.7
Type of closed-loop								
Dual-hormone full closed-loop	5	143	143	19.643 (13.736 to 25.550)	<0.001	0.027	9.41	78.5
Single-hormone full closed-loop	2	245	131	11.671 (8.676 to 14.666)	<0.001		3.45	0.0
Hybrid closed-loop	54	1,904	1,734	10.873 (8.310 to 13.436)	<0.001		87.14	95.0
Type of algorithm								
MPC	37	1,261	1,105	11.411 (8.736 to 14.086)	<0.001	0.915	59.29	88.7
PID	15	595	568	11.595 (7.654 to 15.537)	<0.001		24.09	87.8
Other	10	436	335	13.091 (5.751 to 20.431)	<0.001		16.62	98.6

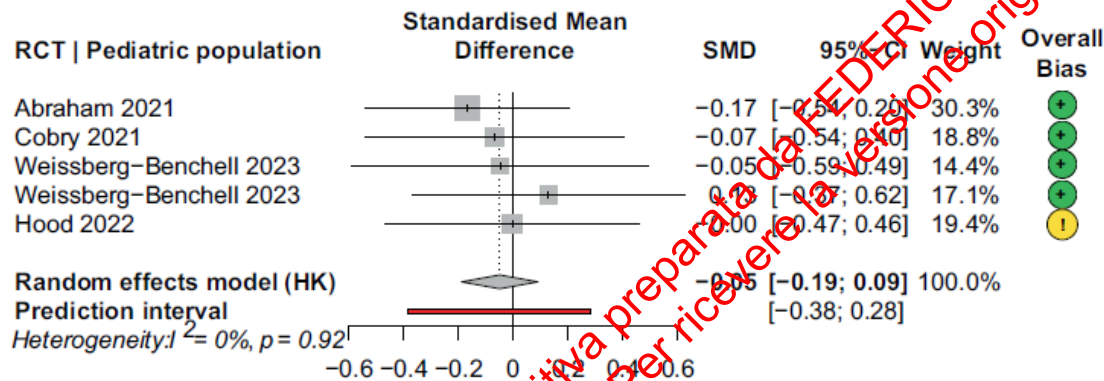
AID & QoL

(a) RCT meta-analysis shows a small beneficial effect of AID on DD in adults

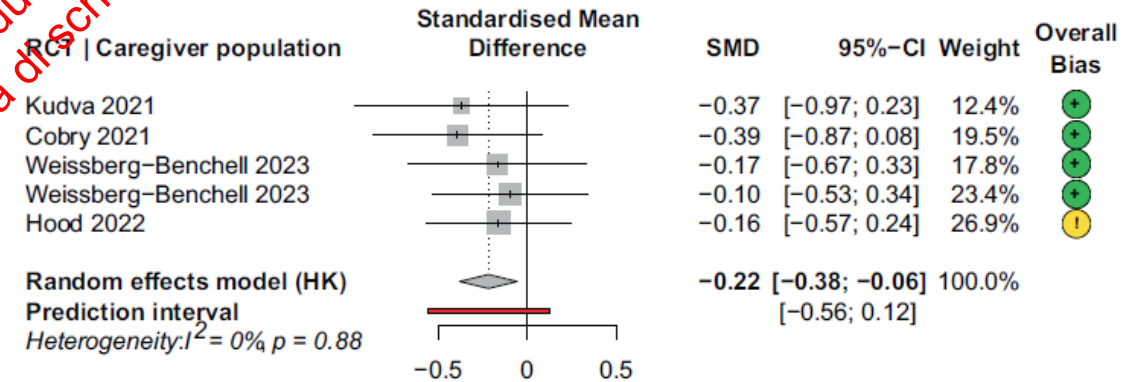
* Only study which included a fully closed-loop system



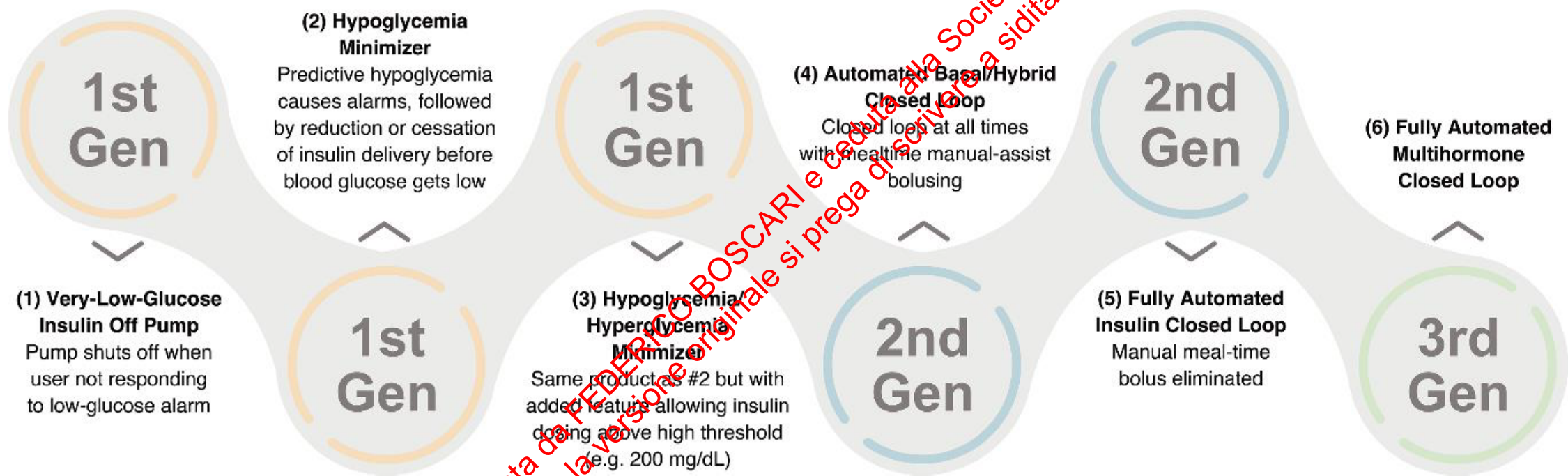
(c) RCT meta-analysis shows no effect of AID on DD in children/teenagers



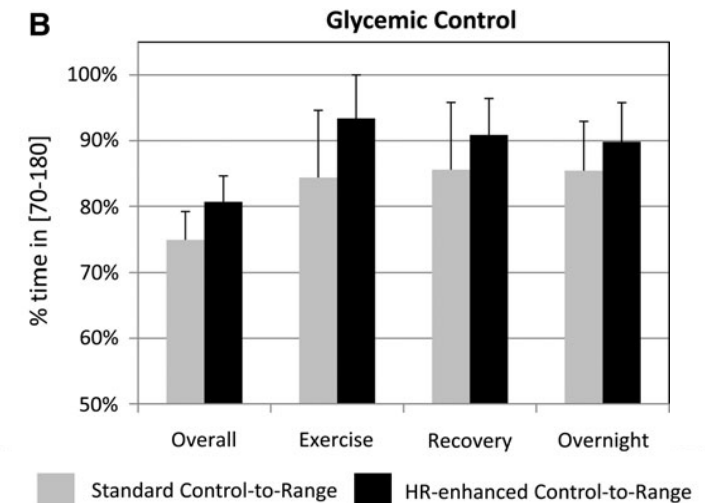
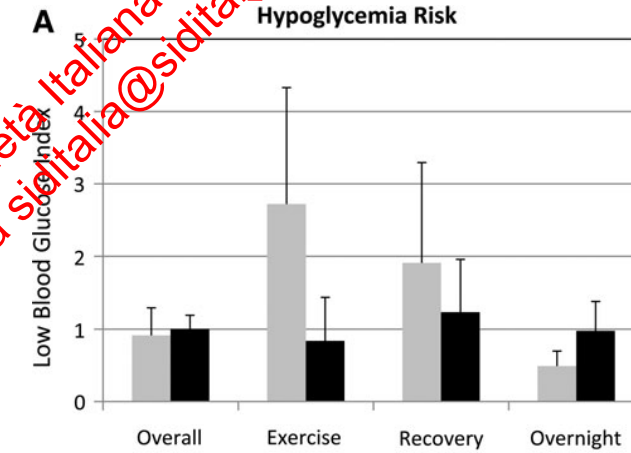
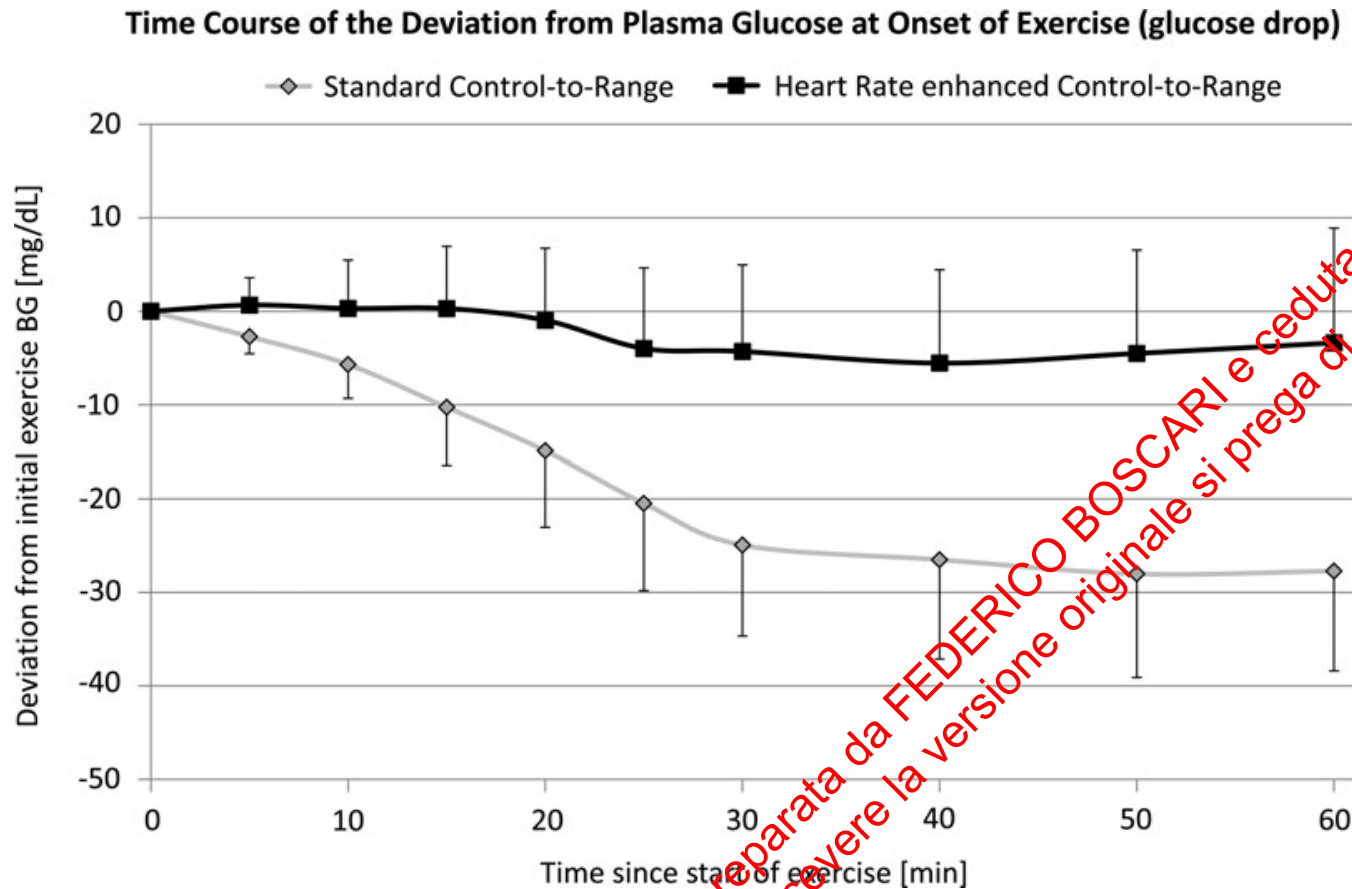
(b) RCT meta-analysis shows a small beneficial effect of AID on DD in caregivers



AID:prospettive future



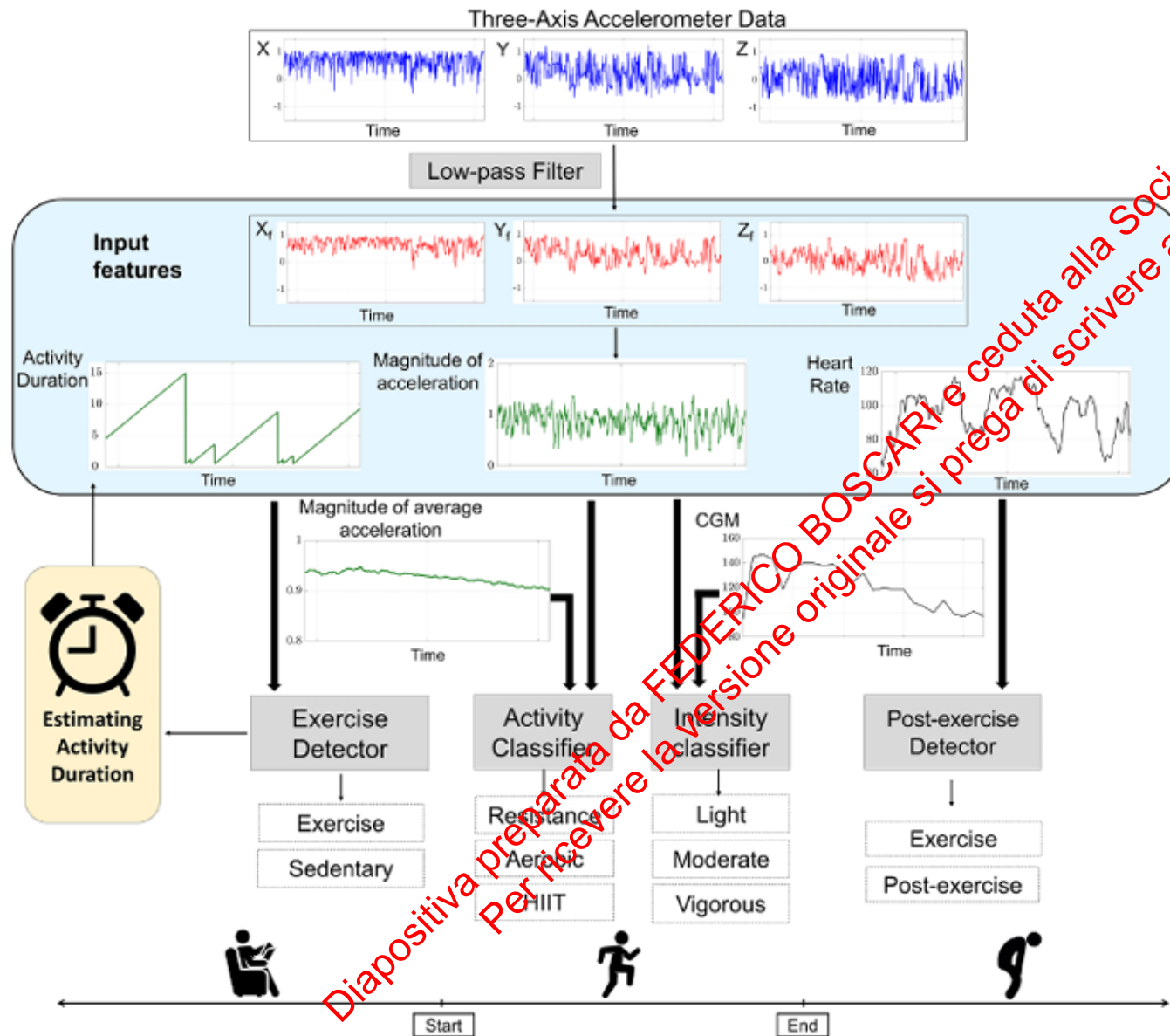
Verso il full closed loop



Breton MD, Brown SA, Karvetskiy CH, Kollar L, Topchyan KA, Anderson SM, Kovatchev BP. Adding heart rate signal to a control-to-range artificial pancreas system improves the protection against hypoglycemia during exercise in type 1 diabetes. Diabetes Technol Ther. 2014 Aug;16(8):506-11. doi: 10.1089/dia.2013.0333. Epub 2014

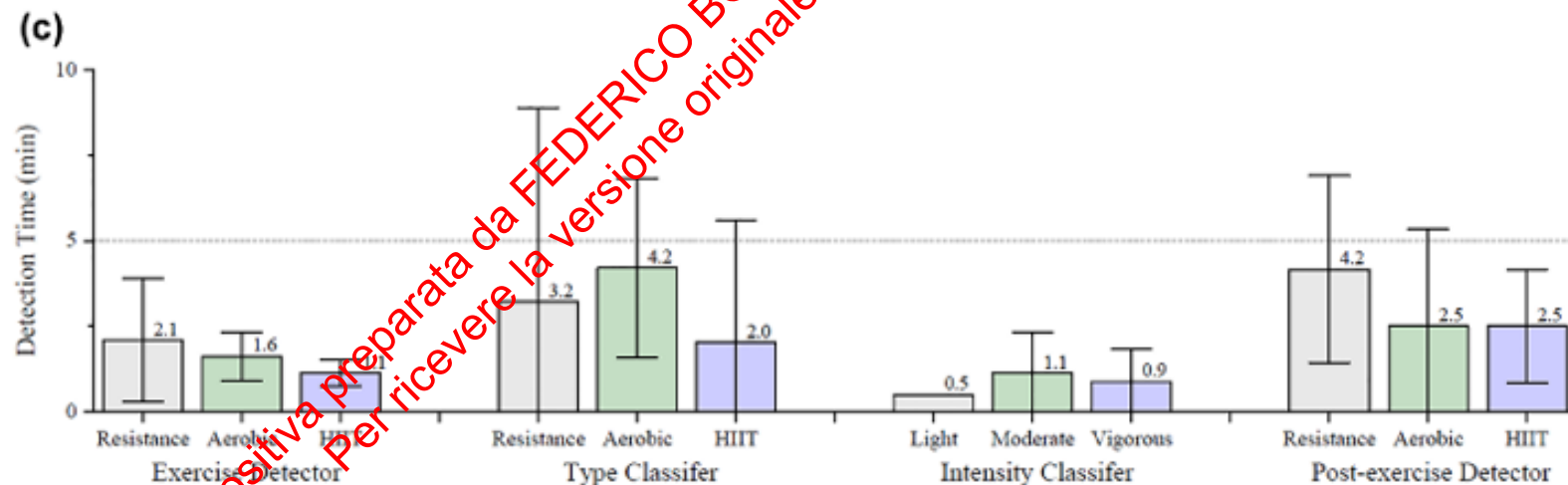
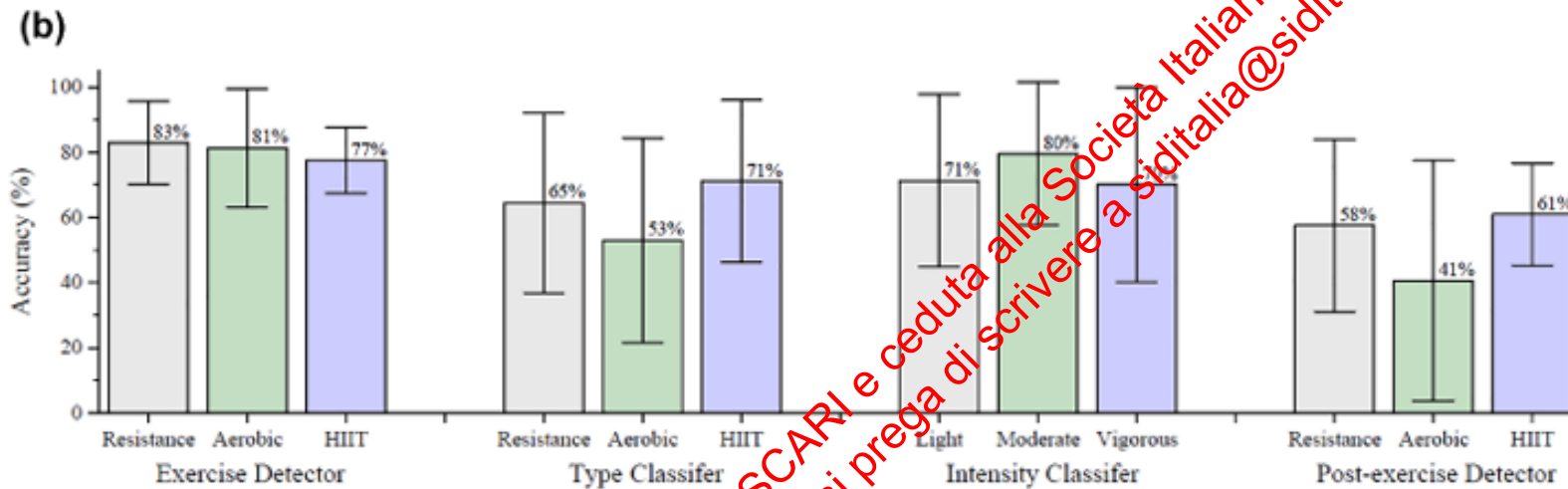
Apr 4. PMID: 24702135; PMCID: PMC4116126.

Come identificare attività fisica



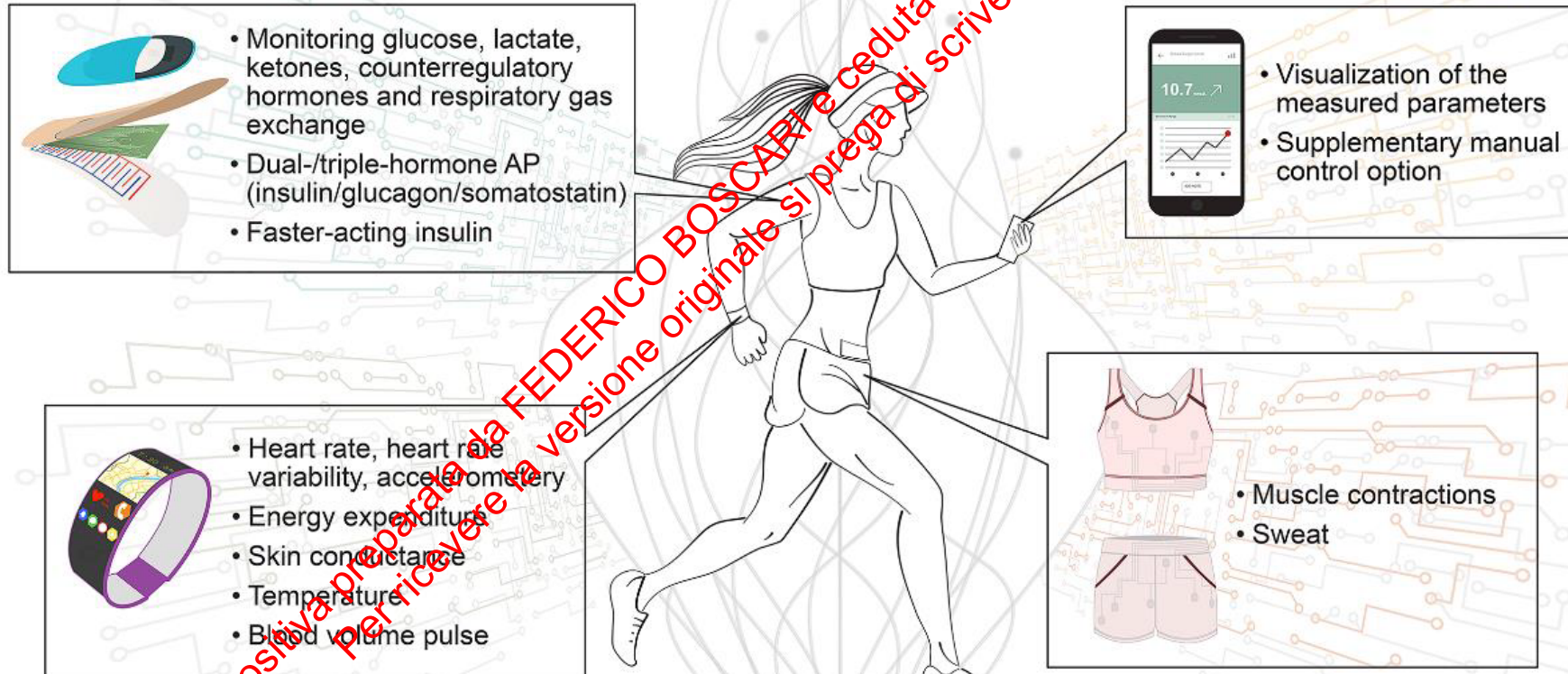
Cho S, Aiello EM, Ozaslan B, Riddell MC, Calhoun P, Gal RL, Doyle FJ 3rd. Design of a Real-Time Physical Activity Detection and Classification Framework for Individuals With Type 1 Diabetes. J Diabetes Sci Technol. 2024 Sep;18(5):1146-1156. doi: 10.1177/19322968231153896. Epub 2023 Feb 17. PMID: 36799284; PMCID: PMC11418461.

Come identificare attività fisica

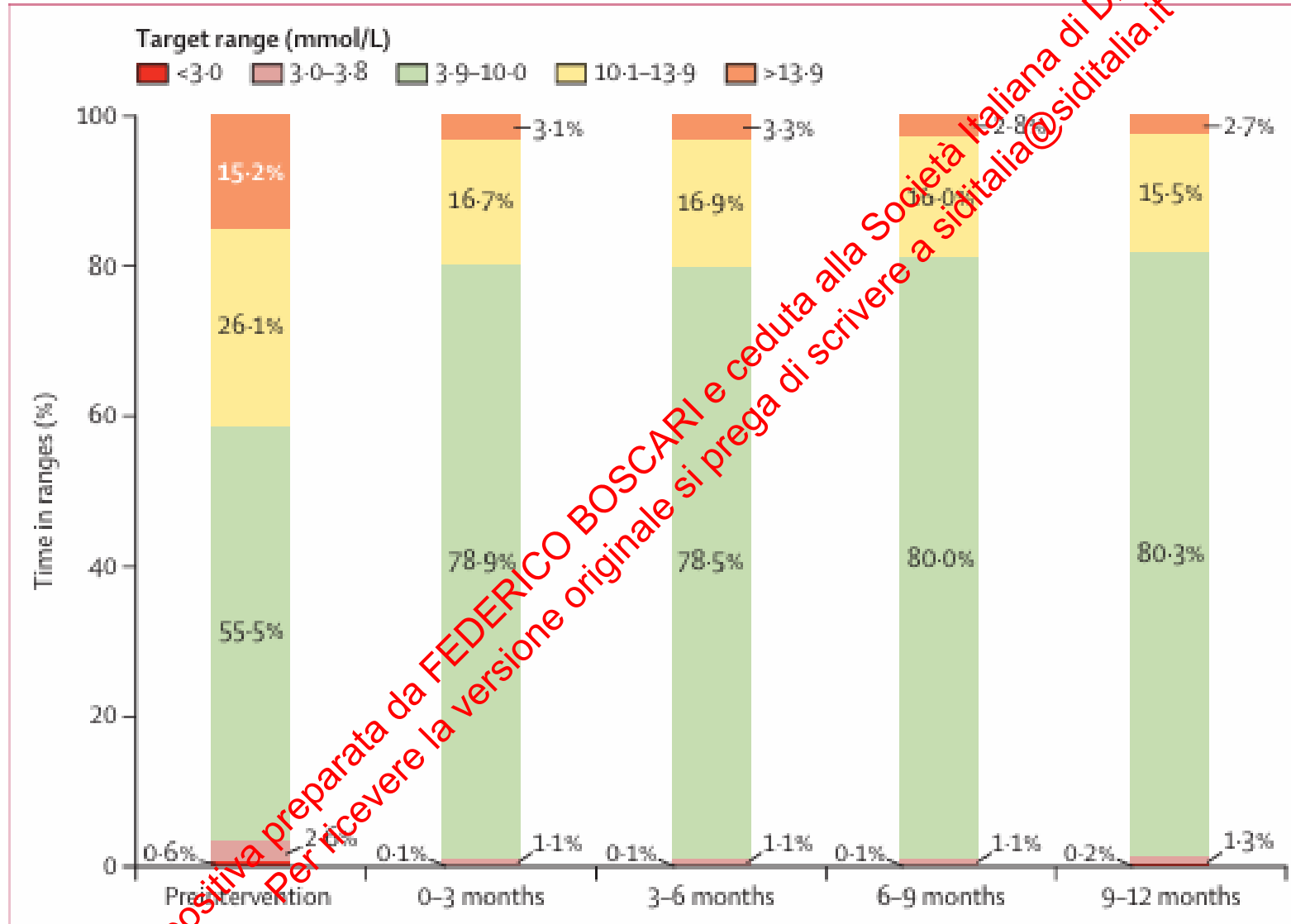


AID:prospettive future

Enhanced algorithms using machine learning and incorporating individual habit patterns and risk profiles



Pancreas biormonale full closed loop



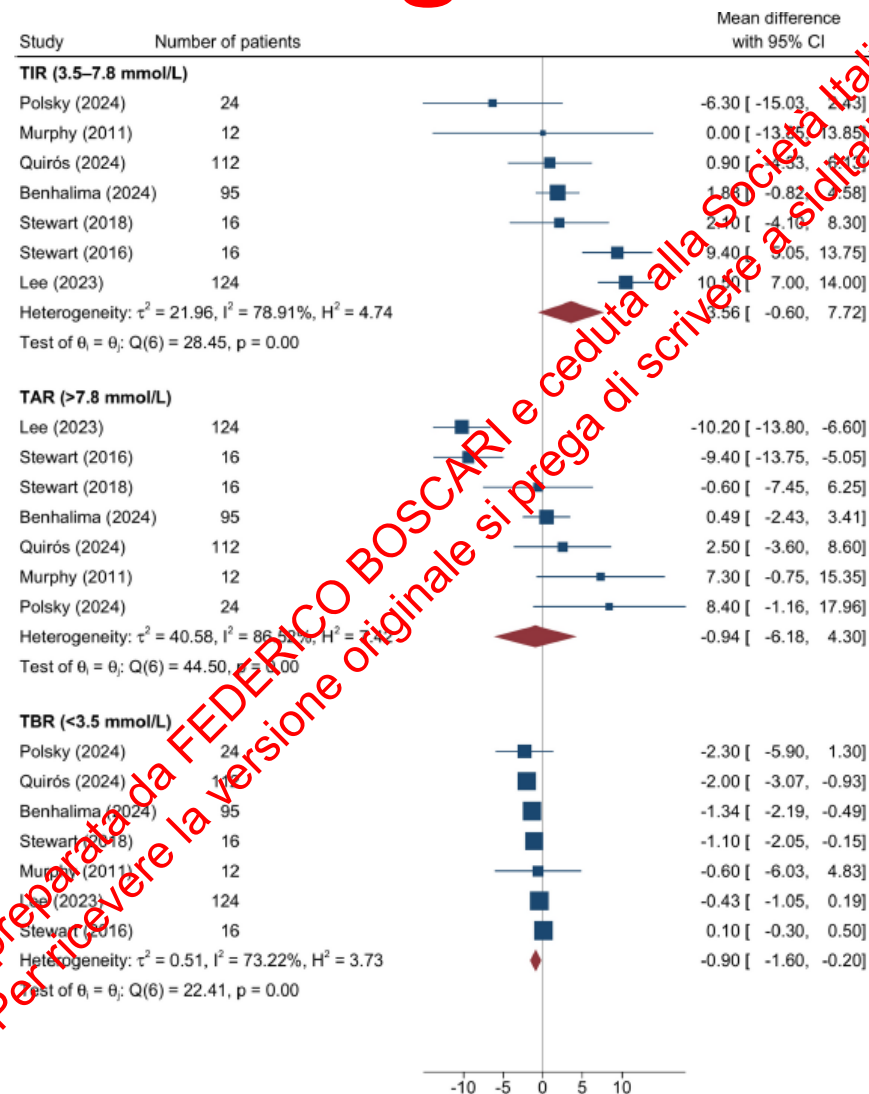
van Bon AC, Blauw H, Jansen TJP, Laverman GD, Urgert T, Geessink-Mennink J, Mulder AH, Out M, Groote Veldman R, Onvlee AJ, Schouwenberg BJW, Vermeulen MAR, Diekman MJM, Gerding MN, van Wijk JPH, Klanssen M, Witkop M, DeVries JH. Bi-hormonal fully closed-loop system for the treatment of type 1 diabetes: a real-world multicentre, prospective, single-arm trial in the Netherlands. Lancet Digit Health. 2024 Apr;6(4):e272-e280. doi: 10.1016/S2589-7500(24)00002-5. Epub 2024 Mar 4. PMID: 38443309.

AID: indicazioni

- DM1
 - gravidanza

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AID in gravidanza



Name study, registration number	Intervention vs comparator	Countries, centers, study end	Study design	Primary outcome(s)	(Main) secondary outcomes	Main results
AiDAPT study (Automated insulin Delivery Amongst Pregnant women with Type 1 diabetes), ISRCTN56898625 ^{29,48}	CamAPS FX AID (Dana Diabecare RS + Dexcom G6 vs CSII or MDI) with Dexcom G6 CGM	UK, 9 centers, Nov 2022	Multicenter, open-label, RCT N = 124 1:1 randomization with stratification by site; Eligibility: GA <14 weeks, HbA1c ≥ 48 mmol/mol ($\geq 6.5\%$) and ≤ 86 mmol/mol ($\leq 10\%$), 18-45 years, T1D diagnosis ≥ 1 year, intensive insulin therapy; Run-in phase of masked Dexcom G6 for baseline data collection	TIRp 3.5-7.8 mmol/L (63-140 mg/dL) between 16 weeks until delivery	HbA1c; DKA; severe hypoglycemic episodes; mode of delivery; GA at delivery; infant birth weight; LGA; SGA; neonatal morbidity; NICU admission; pregnancy loss <24 weeks; stillbirth; neonatal death; fear of hypoglycemia, health status, diabetes distress; sleep quality; cost-effectiveness	AID led to a significant increase in TIRp of 10.5% (mean TIR 68.2% $\pm$ 10.5% in the AID group vs 55.6% $\pm$ 12.5% in the standard of care group), less TARA (difference of 10.2%), higher TIRp overnight (difference of 12.3%), without increase in TBRp and without unanticipated safety problems. Pregnancy outcomes were similar except for a significantly lower rate of hypertensive disorders (20% vs 42%) and less gestational weight gain (11.1 $\pm$ 6.1 vs 14.1 $\pm$ 6.1 kg) in the AID group vs comparator group.
CIRCUIT study (Closed-loop Insulin delivery by glucose Responsive Computer algorithms In Type 1 diabetes pregnancies), NCT04902378 ⁴⁷	Tandem t: slim X2 pump with control IQ with Dexcom G6 CGM vs SoC (MDI or pump) with Dexcom G6	Canada, 9 centers + Australia, 5 centers, 2026	Multicentric open-label RCT N = 94; Eligibility: GA <14 weeks, HbA1c ≥ 48 mmol/mol <86 mmol/mol (≥ 6.5 -<10%), 18-45 years, T1D diagnosis ≥ 1 year, intensive insulin therapy; Run-in phase of one week with Dexcom G6 CGM	TIRp 3.5-7.8 mmol/L (63-140 mg/dL) GA 16-34 weeks gestation	TARp; TBRp; mean blood glucose; Severe hypoglycemia; DKA; fear of hypoglycemia, fear of hyperglycemia; sleep quality; work productivity, preeclampsia; worsening chronic hypertension events; C-sections; preterm delivery; LGA; SGA; mean neonatal birthweight; neonatal hypoglycemia; NICU admission; pregnancy loss	Expected in 2025
CRISTAL study (Closed-loop insulin delivery in pregnant women with type 1 diabetes: a randomized controlled trial), NCT04520977 ^{7,46}	MiniMed 780G system with Guardian 3 or 4 CGM vs continue SoC (MDI or open-loop pump) with any type of sensor	Belgium, 11 centers and the Netherlands 1 center, May 2023	Multicenter, open-label RCT N = 95 1:1 randomization with stratification by center; HbA1c and treatment (pump or MDI) LGA <16 weeks Eligibility: GA <12 weeks, HbA1c ≤ 86 mmol/mol ($\leq 10\%$), 18-45 years, T1D diagnosis ≥ 1 year, intensive insulin therapy; Run-in phase of 10 days with masked Guardian 3 CGM as needed	TIRp 3.5-7.8 mmol/L (63-140 mg/dL) at GA 14-17, 20-23, 26-29, 33-36 weeks	Prespecified: time <3.5 mmol/L (63 mg/dL); overnight time <3.5 mmol/L (63 mg/dL); overnight TIRp 3.5-7.8 mmol/L (63-140 mg/dL) Exploratory: device deficiencies; time >7.8 mmol/L (140 mg/dL); >10 mmol/L (180 mg/dL); time <2.8 mmol/L (50 mg/dL); <3.0 mmol/L (54 mg/dL); severe hypoglycemia; DKA; miscarriage; stillbirth; neonatal death; LGA; respiratory distress; birth trauma; shoulder dystocia; neonatal hypoglycemia; NICU admission; fear of hypoglycemia, health status, diabetes-related emotional distress, treatment satisfaction, cost-effectiveness	The TIRp increased from baseline to the antenatal period [averaged over the four prespecified time points (14–17, 20–23, 26–29, and 33–36 weeks of gestation)] in both groups: from 60.5% $\pm$ 14.2% to 66.5% $\pm$ 10.0% in the AID group and from 57.6% $\pm$ 13.7% to 63.2% $\pm$ 12.4% in the SoC group (adjusted mean difference, 1.88%; 95% CI, -0.82 to 4.58; $P = .17$). Overnight TIRp was higher (adjusted mean difference, 6.58%; 95% CI, 2.31 to 10.85, $P = .0026$), and TBRp overall (adjusted mean difference, -1.34%; 95% CI, -2.19 to -0.49, $P = .0020$) and overnight (adjusted mean difference, -1.86%; 95% CI, -1.86 to -0.81, $P = .0005$) were lower in women with AID compared with the SoC.

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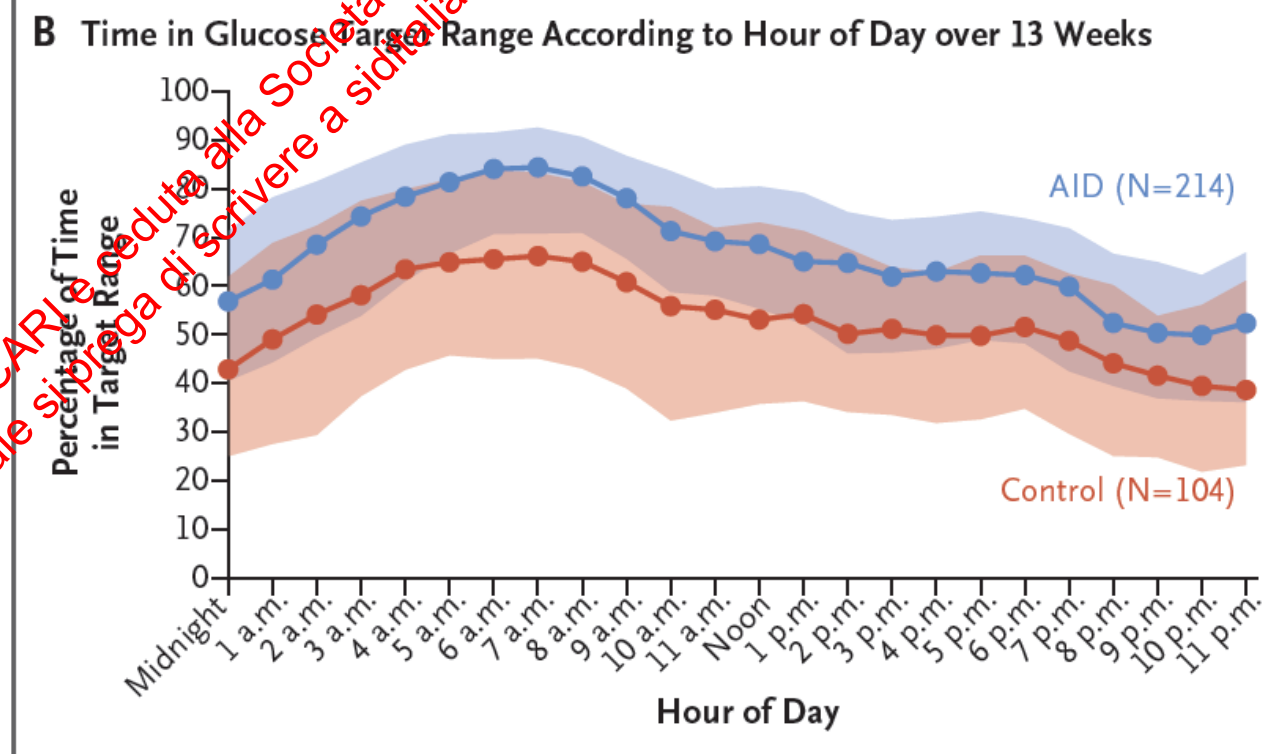
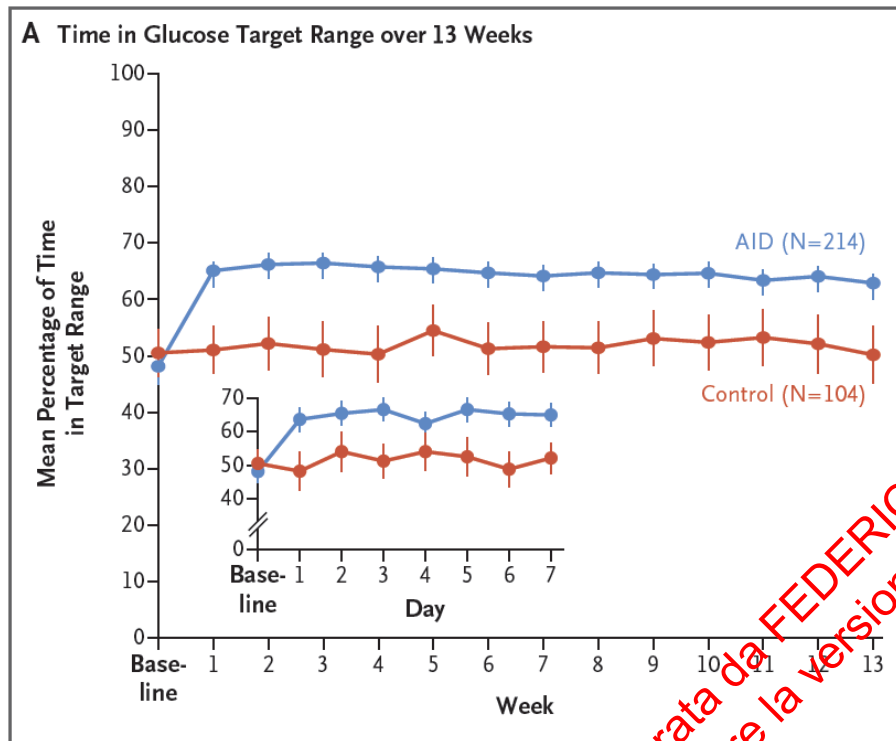
Benhalima K, Polsky S.
Automated Insulin Delivery in
Pregnancies Complicated by
Type 1 Diabetes. J Diabetes
Sci Technol. 2025 Mar
12:19322968251323614. doi:
10.1177/19322968251323614.
Epub ahead of print. PMID:
40071788; PMCID:
PMC11904923.

AID: indicazioni

- DM1
- DM2

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AID nel DM2



Kudva YC, Raghinaru D, Lum JW, Graham TE, Liljenquist D, Spanakis EK, Pasquel FJ, Ahmann A, Ahn DT, Aleppo G, Blevins T, Kruger D, Brown SA, Levy CJ, Weinstock RS, Steenkamp DW, Spaic T, Hirsch IB, Broyles F, Rickels MR, Tsoukas MA, Raskin P, Hatipoglu B, Desjardins D, Terry AN, Singh LG, Davis GM, Schmid C, Kravarusic J, Coyne K, Casaubon L, Espinosa V, Jones JK, Estrada K, Afreen S, Levister C, O'Malley G, Liu SL, Marks S, Peleckis AJ, Pasqua MR, Tardio V, Kurek C, Luker RD, Churchill J, Tajrishi FZ, Dean A, Dennis B, Fronczyk E, Perez J, Mukhashen S, Dhillon J, Ipek A, Bzdick S, Atakov Castillo A, Driscoll M, Averkiou X, Dalton-Bakes CV, Moore A, Jordan LF, Lesniak A, Pinsker JE, Sasson-Katchalski R, Campos T, Spanbauer C, Kanapka L, Kollman C, Beck RW; 2IQP Study Group. A Randomized Trial of Automated Insulin Delivery in Type 2 Diabetes. N Engl J Med. 2025 Mar 19. doi: 10.1056/NEJMoa2415948. Epub ahead of print. PMID: 40105270.

AID: indicazioni

- DM1
- DM2
- Ospedale

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AID in Ospedale

Patient population

Rationale for AID use

Steroid use

Corticosteroid doses are often variable in the hospital and lead to new or rapid changes in insulin requirements

Enteral or parenteral nutrition

Rapid adaptation needed for temporary or unanticipated discontinuation of feeding tubes, cycling of enteral nutrition with oral intake

Type 1 Diabetes

High glycemic variability and hypoglycemia risk

Perioperative period

Acute stress with rapid changes in insulin sensitivity, frequent short term steroid exposure

Pregnancy

Enhanced monitoring and safety are essential for tighter glycemic control during pregnancy to protect both mother and fetus. Insulin needs change throughout pregnancy, sometimes complicated by steroid use for premature deliveries, and dramatically fall after placenta delivery.

Uncontrolled type 2 diabetes

During the acute illness, many patients do not respond to conventional therapy or insulin needs are unknown.

Tecnologia in Ospedale

Recommendations

16.6 In people with diabetes using a personal continuous glucose monitoring (CGM) device, the use of CGM should be continued during hospitalization if clinically appropriate, with confirmatory point-of-care (POC) blood glucose measurements for insulin dosing decisions and hypoglycemia assessment, if resources and training are available, and according to an institutional protocol. **B**

16.7 Continue use of insulin pump or automated insulin delivery in people with diabetes who are hospitalized when clinically appropriate, with confirmatory POC blood glucose measurements for insulin dosing decisions and hypoglycemia assessment and treatment. This is contingent upon availability of necessary supplies, resources, and training, ongoing competency assessments, and implementation of institutional diabetes technology protocols. **C**

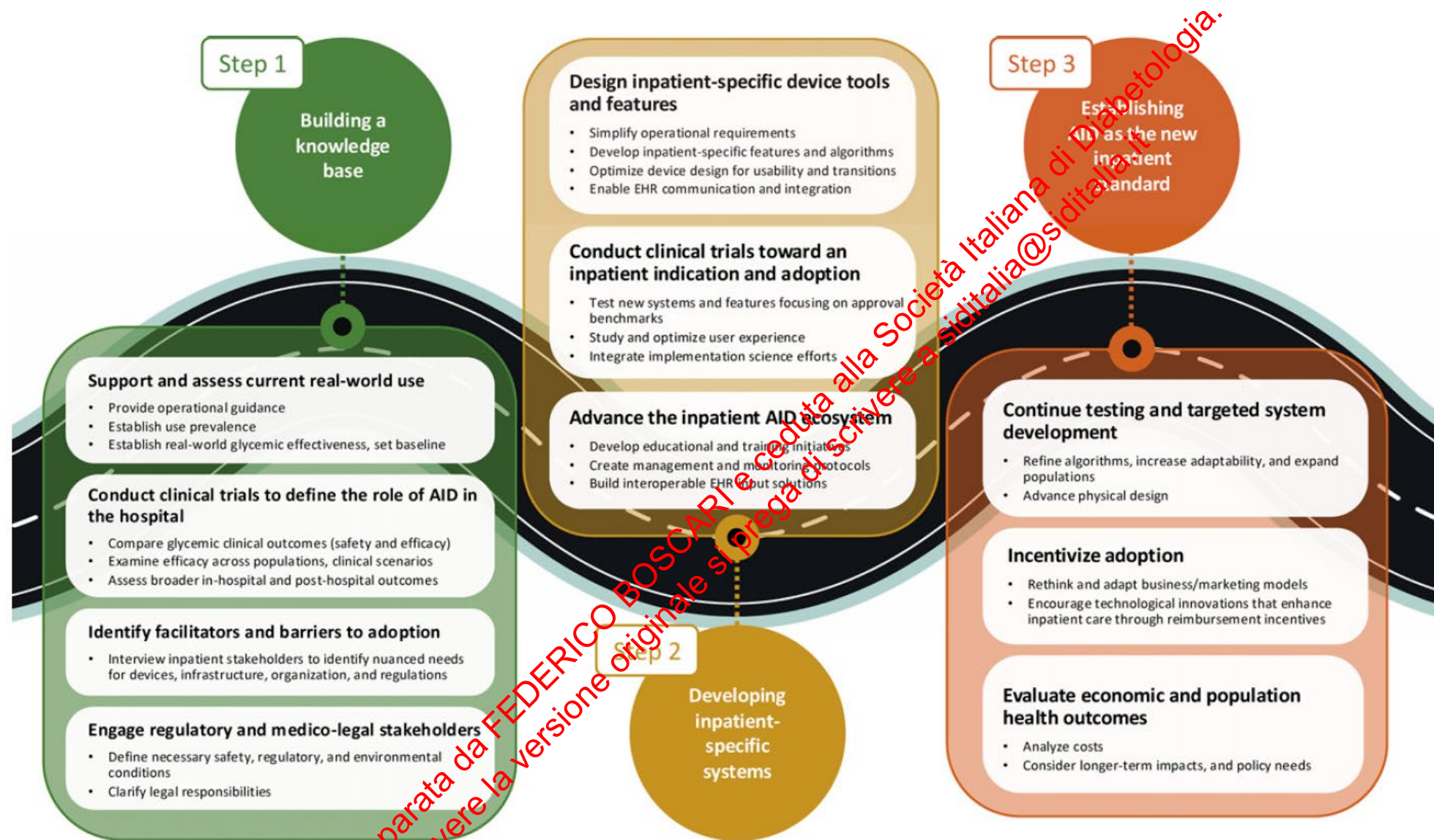


FIG. 1. A roadmap toward making AID an inpatient reality.

Hughes MS, Pasquel FJ, Davis GM, Lal RA, Buckingham BA, Boughton CK, Brown SA, Bally L. Toward Automation: The Road Traveled and Road Ahead for Integrating Automated Insulin Delivery into Inpatient Care. *Diabetes Technol Ther.* 2025 Mar;27(3):217-242. doi: 10.1089/dia.2024.0343. Epub 2024 Dec 2. PMID: 39618315.

Smart MDI



In che modo la tecnologia interviene nella gestione del diabete

Monitoraggio

**Tecnologia &
Diabete**

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Continuous Glucose Monitoring

Lettura sottocutanea
Andamento glicemico
Frecce di tendenza
Allarmi

Similitudini

Peculiarità

Applicazione

Durata

Calibrazioni

Interferenze

Connettività

Accuratezza

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ESSENTIAL FEATURES

(i.e., features that are absolutely necessary for all patients):

- **accuracy**
- **certification**

MAJOR DRIVERS OF CHOICE

(i.e., features that address more relevant clinical issues and/or meet the needs of specific subgroup of patients):

- **approved indications of use**
- **alerts (optional, predictive, on body vibration)**
- **interoperability / integration**
- **approved application sites**
- **visualization devices**
- **transmitter removal**
- **cost of devices**

ADDITIONAL FEATURES

(i.e., features that improve patient's experience):

- **preassembled vs. non-preassembled systems**
- **other alerts (snooze, delay)**
- **device size**
- **notes entry**
- **calibration**
- **allergens**
- **interferences**
- **impermeability**

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Di Molfetta S, Rossi A, Boscari F, Irace C, Laviola L, Bruttomesso D. Criteria for Personalised Choice of a Continuous Glucose Monitoring System: An Expert Opinion. Diabetes Ther. 2024 Nov;15(11):2263-2278. doi: 10.1007/s13300-024-01654-y. Epub 2024 Sep 30. Erratum in: Diabetes Ther. 2025 Jan;16(1):135. doi: 10.1007/s13300-024-01666-8. PMID: 39347900; PMCID: PMC11467157.

La rivoluzione del CGM

- Miglioramento del controllo
- Nuove metriche
- Nuove popolazioni target
- Nuovo ruolo del CGM

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CGM e Obesità

Consensus statements. People living with obesity and at-risk of progression to type 2 diabetes mellitus

- Periodic use of CGM should be considered in people living with obesity and at-risk of developing type 2 diabetes mellitus based on established risk factors,[33] to assess glycemic risks and support early intervention.
- Measures of CGM-defined glycemic risk amongst people living with obesity centred on time above tight range (TATR) >140 mg/dL (7.8 mmol/L) and glycemic variability may be used to identify persons at-risk of intermediate hyperglycemia.
- The application of CGM should be considered during proactive efforts to modify diet and physical activity in people living with obesity, in order to provide immediate feedback on the glycemic impact of dietary choices, as well as the glycemic benefits of physical activity and exercise.

CGM e Prediabete

Consensus statements. People living with intermediate hyperglycemia and at risk of progression to type 2 diabetes mellitus

- Periodic use of CGM should be considered in people living with intermediate hyperglycemia, based on established risk factors for type 2 diabetes mellitus [33] to assess glycemic risks and support early intervention when needed.
- For persons with prior intermediate hyperglycemia and a recent diagnosis of type 2 diabetes mellitus and able to meet CGM targets for >70% TIR, setting targets for TIR can be considered to aim for near-normal glycemia or remission.
- Remission criteria for people being treated for type 2 diabetes mellitus should mirror diagnostic criteria, and include: FPG <126 mg/dL (7.0 mmol/L), mean glucose levels <100 mg/dL (5.6 mmol/L), GMI <6.5% (<48 mmol/mol), sustained over 3 months in the absence of pharmacotherapy.
- Use of CGM should be considered as part of proactive efforts to modify diet and physical activity in people living with intermediate hyperglycemia, in order to provide immediate feedback on the glycemic impact of dietary choices and physical activity or exercise.

Condivisione dato

Caregiver

Dato clinico

Clinica

Telemedicina

Tempo
QoL
Efficacia

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In che modo la tecnologia interviene nella gestione del diabete

Educazione

**Tecnologia &
Diabete**

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Ruolo educativo

1.3 Care systems should facilitate in-person and virtual team-based care, include those knowledgeable and experienced in diabetes management as part of the team, and utilize patient registries, decision support tools, proactive care planning, and community involvement to meet needs of individuals with diabetes. **B**

Improving Care and Promoting Health in Populations. Standards of Care in Diabetes—2025
Diabetes Care 2025;48(Suppl. 1):S14–S260 <https://doi.org/10.2337/dc25-S001>

use of video games, mobile applications, simulators and online interventions, which aim to promote knowledge about DM1, encourage self-care, and improve patients' quality of life. The use of technological resources, which are more attractive and focused on reality, can facilitate training, stimulate reasoning and the ability to solve problems

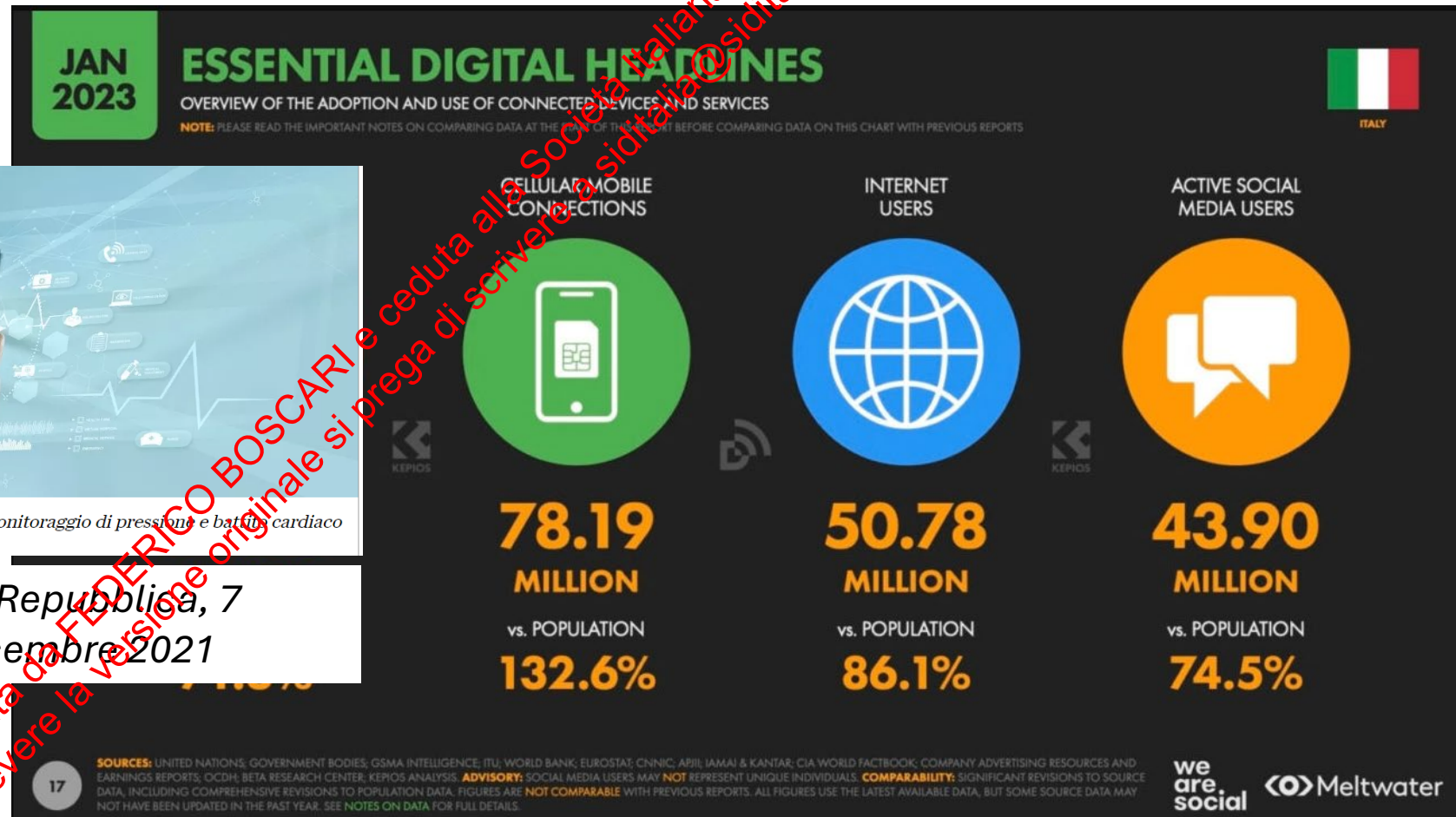
Ruolo educativo

Due italiani su tre utilizzano app e device digitali per controllare il proprio stato di salute

di Salvo Cagnazzo

Le app più gettonate sono quelle dedicate a dieta e alimentazione (28%), al monitoraggio di pressione e battito cardiaco (24%), alla regolarità del ciclo mestruale (20%)

La Repubblica, 7
Dicembre 2021



APP & Diabete

- Registrare dati clinici
- Ruolo nel CHO Counting e calcolo del bolo
- Condivisione dati
- Ruolo nell'adottare stili di vita salutari
- Migliorare conoscenza diabete
- Prevenire complicanze
- Interazione

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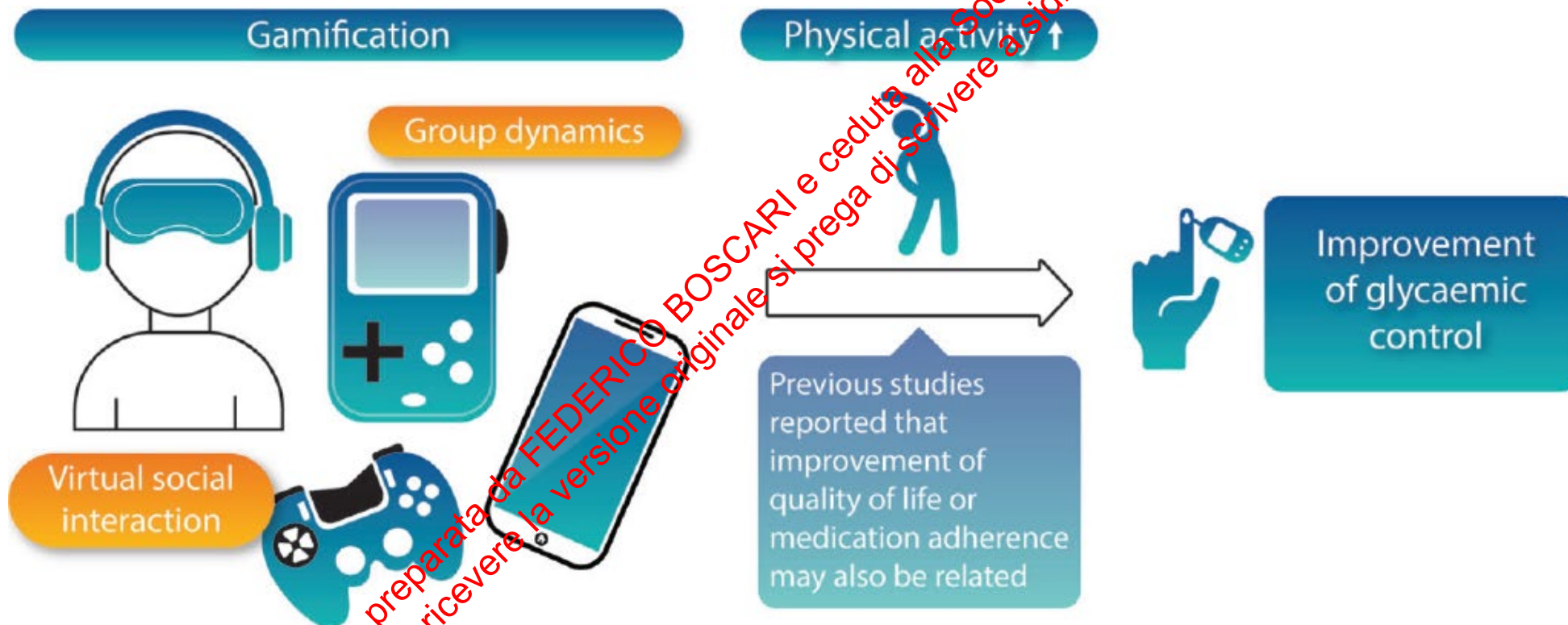
APP & CHO Counting

Table 2. Carbohydrate counting and glycemic control outcomes (at baseline and follow-up).

Outcomes	Baseline			Follow-up			Coefficient
	iSpy (n=22)	Usual care (n=22)	P value	iSpy (n=21)	Usual care (n=22)	P value	
Absolute error (% of total grams), mean (SD)	31.97 (11.36)	32.08 (10.01)	.99	27.45 (10.90)	38.00 (14.74)	.008	−10.479
Errors >10 g (%), mean (SD)	25.00 (14.06)	27.73 (15.10)	.54	21.43 (16.82)	32.27 (16.31)	.047	−9.851
Total time (seconds), mean (SD)	79.95 (23.88)	80.73 (27.82)	.92	74.86 (31.78)	78.23 (44.97)	.79	−2.741
HbA _{1c} ^a (%), mean (SD)	8.41 (1.84)	8.35 (1.32)	.91	8.06 (1.43)	8.80 (1.60)	.03	−0.603

Alfonsi JE, Choi EEY, Arshad T, Sammott SS, Pais V, Nguyen C, Maguire BR, Stinson JN, Palmert MR. Carbohydrate Counting App Using Image Recognition for Youth With Type 1 Diabetes: Pilot Randomized Control Trial. JMIR Mhealth Uhealth. 2020 Oct 28;8(10):e22074. doi: 10.2196/22074. PMID: 33112249; PMCID: PMC7657721.

APP & Diabete



Name of the Game	Description of the Game	Features of the Game	Game Themes	Game Mechanics	Operating Systems
Gluko Bear	The game aims to provide education regarding the best nutrition for diabetic patients. It involves a bear which get through different levels by matching healthy food and keeping the blood glucose level low.	- Available in three languages. (English, German, Portuguese) - Includes quizzes - More than 100 types of food with different sugar levels. - Targeted age group +3	Quizzes	Levels	Android
Pandabetic	It aims to educate type 1 diabetic children through entertainment. The children need to maintain the glucose level of the character through managing insulin and carbohydrates.	- Available in three languages: (English, Spanish, Italian) - Social interaction through the support of game center. (for iOS) - It supports family sharing feature for up to six family members. (for iOS) - Targeted age group +3	Taking care of a character	Levels and points	Both Android and iOS
Diapets	The game encourages the diabetic child to check the blood levels on time, help to overcome the fear of insulin needles and to count carbs. This is done through taking care of a baby dragon character.	Notification feature for reminding for the eating time and for using blood glucose meter. It supports family sharing feature for up to six family members. Targeted age group +4, 6–8	Taking care of a character	Points	iOS
GlukoZor World	The game aims to provide education about diabetes in general by taking care of the game character. The user can choose the color and the weight of the character. The user can feed the character, entertain him by many activities that need to be unlocked, and managing his blood glucose level.	- Available in four languages. (English, Dutch, Italian, Spanish) - It includes quizzes - It supports family sharing feature for up to six family members. - Targeted age group +3	Taking care of a character	Levels, points and dashboard	Both Android and iOS
Diabetes control	The game is generally a glycemic index which includes certain type of food and their amount of sugar. The user shall choose among two type of food which include less amount of sugar.	- More than 30 comparison of more than 60 types of food - Great to memorize - Targeted age group +3	Quizzes	N/A	Android

Alsalman D, Bu Ali ZM, Alnosaier ZF, Alotaibi NA, Alanzi TM. Gamification for Diabetes Type 1 Management: A Review of the Features of Free Apps in Google Play and App Stores. J Multidiscip Healthc. 2020 May 14;13:425-432. doi: 10.2147/JMDH.S249664. PMID: 32523349; PMCID: PMC7236238.

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and meta-analysis of randomized
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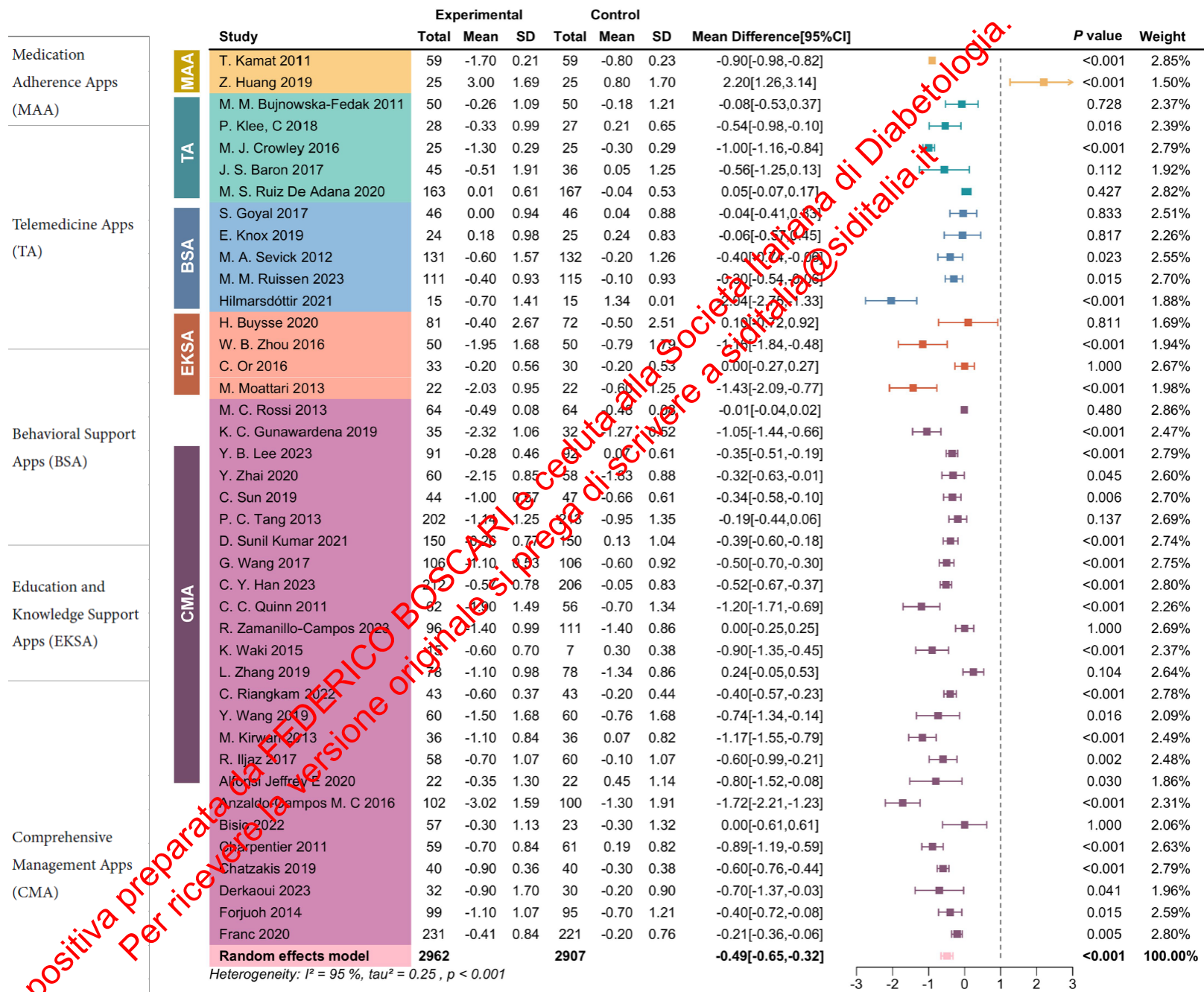


FIGURE 2
Forest plot illustrating the effect of mobile health on HbA1c.

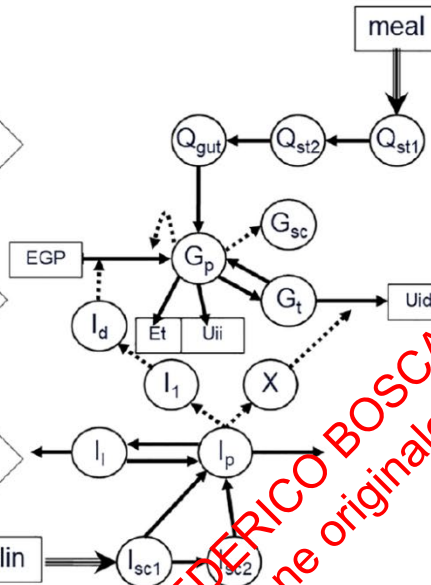
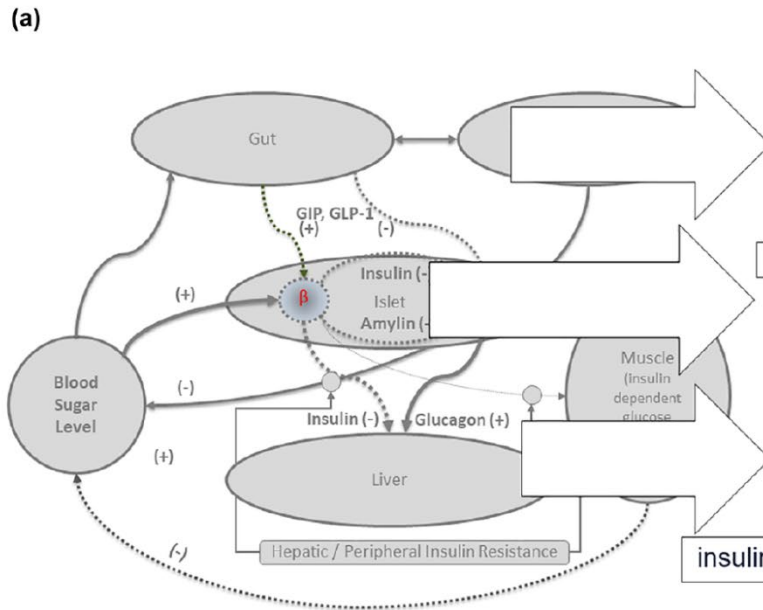
In che modo la tecnologia interviene nella gestione del diabete

**Tecnologia &
Diabete**

Ricerca

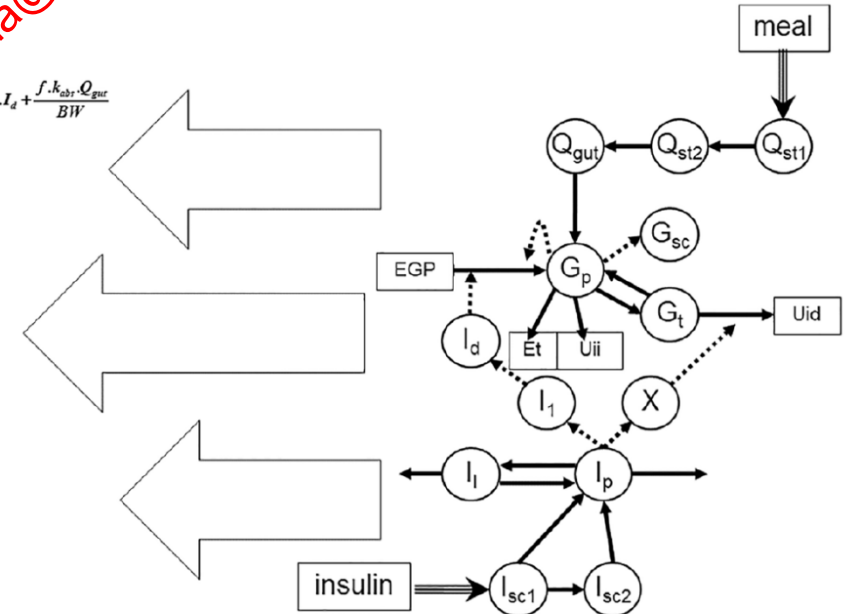
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Tecnologia & Ricerca

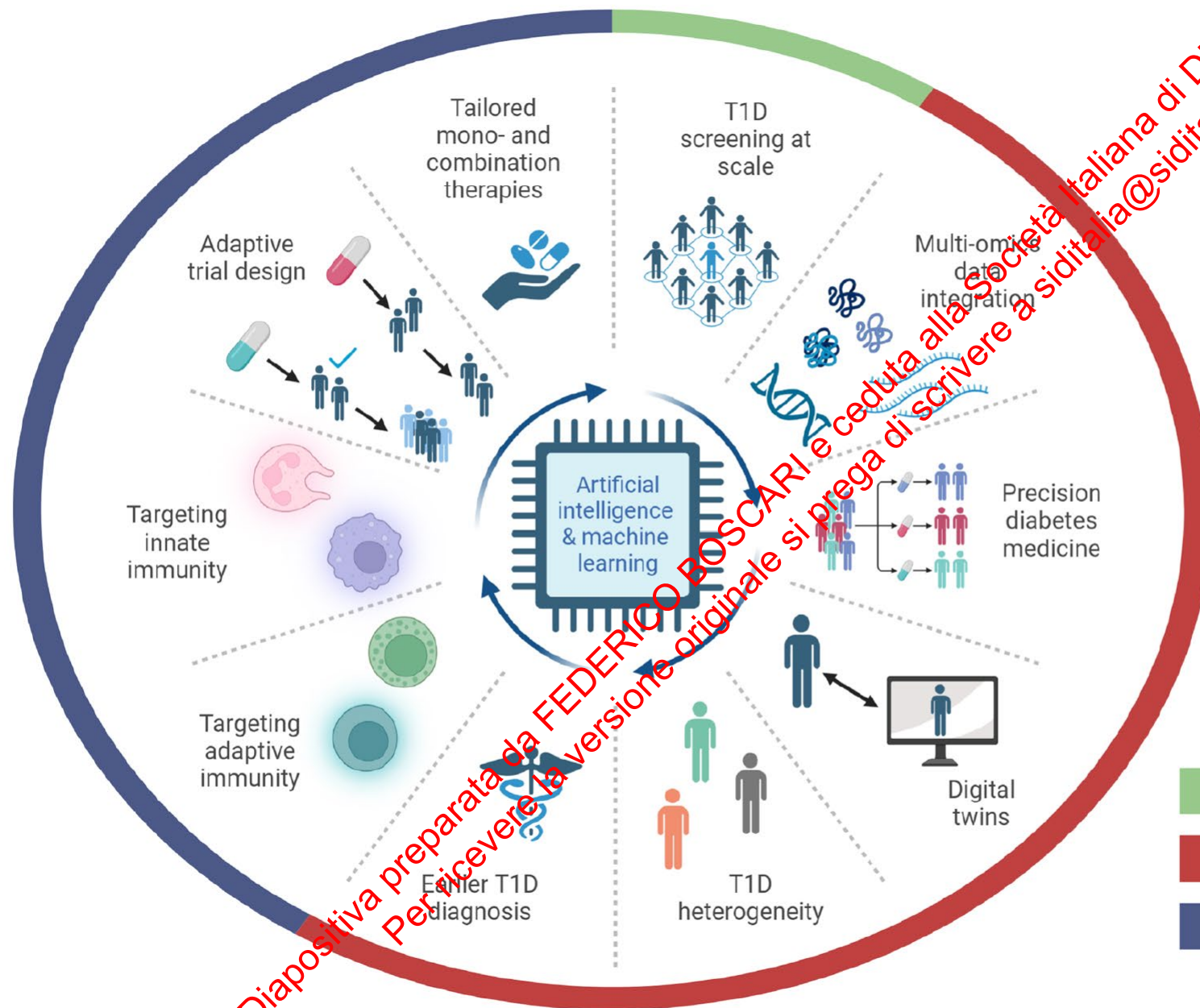


(b)

$$\begin{aligned} \dot{G}_p &= -k_1 \cdot G_p + k_1 \cdot G_t - I_p \cdot G_p + k_{p1} \cdot G_p - k_{p2} \cdot I_d + \frac{f \cdot k_{abr} \cdot Q_{gut}}{BW} \\ \dot{G}_t &= -k_1 \cdot G_t + k_1 \cdot G_p - \frac{(V_{m0} + V_{m1} \cdot G_p)}{K_{m0} + G_p} \cdot G_t \\ \dot{G}_{sc} &= -k_1 \cdot G_{sc} + \frac{G_p}{V_{sc}} \\ \dot{I}_p &= m_1 \cdot G_p + m_1 \cdot G_{sc} + m_1 \cdot I_i + k_{a1} \cdot I_{sc1} + k_{a2} \cdot I_{sc2} \\ &\quad - (m_2 + m_3) \cdot I_i + m_2 \cdot I_p \\ \dot{I}_1 &= k_1 \cdot \left(I_d - \frac{I_p}{V_i} \right) \\ \dot{I}_d &= -k_1 \cdot (I_d - I_1) \\ \dot{X} &= -p_{2a} \cdot \left(X - \left(\frac{I_p}{V_i} - I_d \right) \right) \\ \dot{I}_{sc1} &= -k_d \cdot I_{sc1} - k_{a1} \cdot I_{sc1} + \frac{J(t)}{BW} \\ \dot{I}_{sc2} &= k_d \cdot I_{sc1} - k_{a2} \cdot I_{sc2} \\ \dot{Q}_{gut1} &= -k_{gut1} \cdot Q_{gut1} + M(t) \\ \dot{Q}_{gut2} &= -k_{gut2} \cdot Q_{gut2} + k_{gut1} \cdot Q_{gut1} \\ \dot{Q}_{gut} &= k_{abr} \cdot Q_{gut} + k_{emp1} \cdot Q_{gut2} \end{aligned}$$



Cobelli C, Kovatchev B. Developing the UVA/Padova Type 1 Diabetes Simulator: Modeling, Validation, Refinements, and Utility. J Diabetes Sci Technol. 2023 Nov;17(6):1493-1505. doi: 10.1177/19322968231195081. Epub 2023 Sep 25. PMID: 37743740; PMCID: PMC10658679.



Shapiro MR, Tallon EM, Brown ME, Posgai AL, Clements MA, Brusko TM. Leveraging artificial intelligence and machine learning to accelerate discovery of disease-modifying therapies in type 1 diabetes. *Diabetologia*. 2025 Mar;68(3):477-494. doi: 10.1007/s00125-024-06339-6. Epub 2024 Dec 19. PMID: 39694914; PMCID: PMC11832708.

- Cohort selection
- Cohort, therapeutic agent and endpoint selection
- Therapeutic agent and endpoint selection

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I limiti della tecnologia

- Educazione
- Formazione paziente/clinico
- Diffusione
- Equità

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Table 1. Prevalence of type 1 diabetes and usage of diabetes technology by selected countries and regions^a

	Overall prevalence of diabetes (percentage of overall population with diabetes)	Percentage of overall population with diabetes that have type 1 diabetes	Percentage of patients with type 1 diabetes that use medical technology (pump/CSII, CGM, or closed-loop hybrid system/automated insulin delivery system)	Percentage of adolescent and young adult patients with type 1 diabetes that use medical technology (pump/CSII, CGM, or closed-loop hybrid system/automated insulin delivery system)			
North America							
Canada	5.88 million 15% of population [10]	5%-10% [11]	35% pump 24% CGM [12]	No data		Conway RB, Snell-Bergeon J, Honda-Konno K, Peddi AK, Isa SB, Sulong S, Sibomana L, Gerard Gonzalez A, Song J, Lomax KE, Lo CN, Kim W, Haynes A, de Bock M, Burckhardt MA, Schwab S, Hong K. Disparities in Diabetes Technology Uptake in Youth and Young Adults With Type 1 Diabetes: A Global Perspective. J Endocr Soc. 2024 Nov 28;9(1):bvae210. doi: 10.1210/jendso/bvae210. PMID: 39703363; PMCID: PMC11655873.	
United States	38.4 million 11.6% of population [13]	5%-10% [13]	66% CGM [14] 41% pump [14]	40% CGM [14]			
Latin America							
Mexico	14.1 million 14.1% of population	6% [15]	21% CSII [15] 9% CGM	No data			
Asia							
Korea	4.4 million 13.9% of population	1.1%	10.7% CGM [16, 17]	14.0% pump 39.3% CGM [18]			
Japan	11.96 million 14.6% of population [19]	2.4% [20, 21]	39% pump 11% SAP 48.4% CGM [22]	21%-28% SAP [22]			
Taiwan	2.186 million 11.3% of adults [23]	0.24%	No data	No data			
Europe/Australia/New Zealand							
Malaysia	3.6 million 19% of population [24]	0.62% [25]	<10% [26]	Australia	4.3 million 5.3% population [31]	9.6% [31]	33.3% pump 80.3% CGM
India	101 million 11.4% of population [25]	5%-10%	<10% pump No data on CGM	New Zealand	268 700 6.2% of population	5-10% [33]	No data
Africa							
Rwanda	400 000 [1] 4.5% of population	0.9% [1]	Technology not available				
Kenya	460 000 [27]-821 500 [28] 4.2% [27] to 5.1% [28] of population	≤10% [27]	3% CGM [29]	Switzerland	400 000-500 000 of 8.74 million (~5% of the population) [28, 34]	0.8%-1% [1]	No data
South Africa	4.6 million [30] 10.5% [30] to 11.3% of population [28]	7% [1, 28]	1% pump 16% CGM [28]	Other Europe	61 million 7% of population	5%	Germany: 57% pump 49% CGM [35] Denmark: 39% pump 60% CGM [36] Norway: 74% pump 52% CGM [37] Sweden: 70% pump (children) [38] and 30% pump (adults) [39] 80% CGM [38]

Conclusioni

- La tecnologia ha un ruolo fondamentale nella gestione complessiva del diabete
- Ruolo dei clinici e delle società scientifiche deve essere quello di garantire un accesso equo a tali risorse e di formare il personale alla corretta gestione delle tecnologie stesse

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