

1 Comprehensive Diabetic Care Program Reduces Glycemic 2 Burden and Cardiovascular Risk Markers in Type 2 Diabetes: 3 A Retrospective Cohort Study 4

5 ABSTRACT

6 **Background:** Type 2 diabetes mellitus (T2DM) is a leading driver of cardiovascular morbidity
7 globally, with conventional pharmacotherapy offering glycemic control but limited benefit on the
8 broader cardiometabolic risk profile. Integrative Ayurvedic programs combining dietary
9 restriction, Panchakarma procedures, and structured exercise offer a multimodal approach that
10 may address glycaemia and cardiovascular risk factors simultaneously. Evidence from rural and
11 semi-urban Indian populations, where environmental and lifestyle determinants differ
12 substantially from urban cohorts, remains limited.

13 **Objective:** To evaluate the impact of the Comprehensive Diabetic Care (CDC) program on
14 glycemic parameters, anthropometric indices, blood pressure, resting heart rate, and
15 antidiabetic medication burden in patients with T2DM at a semi-rural clinic in the Marathwada
16 region of Maharashtra, India.

17 **Methods:** A retrospective pre-post cohort study of 32 T2DM patients who completed the CDC
18 program (mean 10.3 ± 3.6 Panchakarma sessions). The program comprised caloric restriction
19 (~ 800 kcal/day; Prameha diet), Panchakarma procedures (Neem Siddha Taila Abhyanga,
20 DashmulaKwath Swedan, and Kwath-based Basti delivering berberine, gymnemic acids, and
21 glycyrrhizin), and yoga-based exercise. Paired t-tests assessed pre-post differences; Pearson
22 correlation evaluated the dose-intensity relationship between Panchakarma sessions and
23 outcome variables.

24 **Results:** The program produced significant improvements across all cardiometabolic domains.
25 HbA1c fell from $8.67 \pm 2.28\%$ to $7.47 \pm 1.68\%$ ($\Delta -1.19\%$, 95% CI -0.67 to -1.71 , $p < 0.001$).
26 Body weight declined by 4.28 kg ($p < 0.0001$), BMI by 1.52 kg/m² ($p < 0.0001$), and abdominal
27 girth by 4.73 cm ($p < 0.0001$). Systolic blood pressure fell by 7.16 mmHg ($p = 0.011$) and
28 diastolic by 6.45 mmHg ($p = 0.006$). Resting heart rate decreased by 7.68 bpm ($p < 0.001$).
29 Fasting blood glucose declined by 59.9 mg/dL ($p = 0.001$). Fifty percent of participants achieved
30 HbA1c $< 7.0\%$ and 37.5% achieved HbA1c $< 6.5\%$. Complete antidiabetic medication

31 discontinuation was achieved in 11 of 28 medicated patients (39.3%). Panchakarma dose
32 intensity significantly correlated with weight reduction ($r = -0.46$, $p = 0.008$).

33 **Conclusion:** The CDC program produced clinically meaningful and statistically significant
34 improvements across glycemic, anthropometric, hemodynamic, and autonomic parameters in a
35 semi-rural T2DM cohort, with a dose-dependent effect on adiposity. These findings support the
36 role of structured multimodal Ayurvedic intervention as a comprehensive cardiometabolic risk
37 reduction strategy in T2DM.

38 **Keywords:** Madhavbaug, T2DM; Ayurveda; Panchakarma; glycemic control; cardiovascular risk;
39 blood pressure; heart rate; Prameha diet; berberine; Gymnemic acid, Sangamner

40 1. INTRODUCTION

41 Type 2 diabetes mellitus (T2DM) has reached epidemic proportions in South Asia, with India
42 hosting an estimated 101 million people with the condition — a figure projected to rise
43 substantially by 2045.^{1,2} The disease burden extends far beyond hyperglycemia: T2DM
44 substantially elevates the risk of cardiovascular disease, hypertension, and obesity-related
45 complications, which together account for the majority of T2DM-related mortality and disability.³

46 Conventional pharmacotherapy achieves glycemic targets in a proportion of patients but has
47 limited primary effect on cardiovascular risk factors beyond glycaemia, and is associated with
48 long-term medication burden, weight gain with certain agents, and suboptimal adherence in
49 resource-limited settings.^{4,5} There is thus growing interest in integrative and lifestyle-based
50 interventions that address the broader cardiometabolic risk profile of T2DM, not glycaemia
51 alone.

52 Ayurvedic medicine conceptualizes T2DM under the framework of Prameha — a disorder
53 characterized by excess of Kapha and Medas (adipose tissue) — and prescribes multimodal
54 intervention combining dietary modification, purificatory Panchakarma procedures, and
55 exercise.^{6,7} The Comprehensive Diabetic Care (CDC) program operationalizes this framework as
56 a structured clinical protocol delivering caloric restriction through a low-carbohydrate Prameha
57 diet, Panchakarma comprising Abhyanga (oil massage), Swedan (medicated steam), and
58 Kwath-based Basti (enema delivering pharmacologically active phytochemicals), and structured
59 yoga and exercise targeting muscle insulin sensitivity.

60 Prior studies from urban clinic cohorts have demonstrated the efficacy of the CDC program in
61 reducing HbA1c, body weight, and antidiabetic medication burden.^{8,9} However, the simultaneous
62 effect of this program on multiple cardiovascular risk domains — including blood pressure and

63 resting heart rate — has not been comprehensively characterized, particularly in semi-rural
64 populations where lifestyle, dietary patterns, and comorbidity profiles differ from urban settings.
65 The present study evaluates the effect of the CDC program on glycemic parameters,
66 anthropometric indices, blood pressure, resting heart rate, and antidiabetic medication burden in
67 a cohort of T2DM patients attending a semi-rural clinic in the Marathwada region of
68 Maharashtra, India. We hypothesized that the program would produce significant improvements
69 across all cardiometabolic domains simultaneously, consistent with its multimodal mechanism of
70 action.

71 **2. METHODS**

72 **2.1 Study Design and Setting**

73 This was a retrospective pre-post cohort study conducted at a semi-rural Madhavbaug clinic in
74 the Marathwada region (Ahmednagar district), Maharashtra, India. Patients enrolled in the CDC
75 program between April 2025 and March 2026 were included. The Marathwada region is
76 characterized by an agrarian economy, high rates of T2DM, and limited access to specialist
77 diabetology services, making it a representative population for evaluating accessible integrative
78 interventions.

79 **2.2 Participants**

80 Inclusion criteria: adults with a confirmed diagnosis of T2DM, who enrolled in the CDC program
81 and had complete pre- and post-intervention HbA1c data. Patients with primary diagnoses of
82 non-diabetic obesity, dyslipidemia, or hypertension without T2DM were excluded. Of 38 records
83 retrieved, 32 met inclusion criteria and constituted the analytic cohort.

84 **2.3 The CDC Program**

85 The CDC program is a structured multimodal intervention delivered in tiered packages (CDC
86 SP1–SP3 and CDC KP1–KP2) corresponding to 8–20 planned Panchakarma sessions across a
87 five-month active treatment period, followed by monthly follow-up. Sessions were administered
88 once weekly for the initial three months (intensive phase), twice monthly for the subsequent two
89 months (consolidation phase), and once monthly thereafter (maintenance follow-up). The
90 program comprises three integrated components:

91 **Dietary Intervention:** Caloric restriction to approximately 800 kcal/day using the Prameha diet
92 — a low-carbohydrate diet emphasizing complex polysaccharides, Millets, Par boiled rice,
93 Barley, pulses, fenugreek, and elimination of refined sugars and high-glycemic-index foods,
94 consistent with current nutritional guidelines for T2DM management.

95 **Panchakarma Procedures:** A sequential protocol comprising three procedural steps delivered
96 at each session: (i) Centripetal oleation (Abhyanga) — whole-body massage with Neem Siddha
97 Taila administered in centripetal strokes, facilitating transdermal delivery of anti-inflammatory
98 and insulin-sensitizing phytochemicals and enhancing peripheral circulation; (ii) Thermal
99 vasodilation (Swedan) — medicated steam exposure using DashmulaKwath decoction,
100 producing controlled hyperthermia and vasodilation that augments metabolic activation and
101 phytochemical tissue penetration; and (iii) Per-rectal drug administration (PRDA / Basti) — a
102 retention enema delivering a Kwath-based decoction containing gymnemic acids (intestinal
103 glucose transport inhibitors derived from *Gymnemasylvestre*), glycyrrhizin (anti-inflammatory
104 saponin derived from *Glycyrrhiza glabra*), and curcuminoids (NF- κ B inhibitors and insulin
105 sensitizers derived from *Curcuma longa*) via the colono-portal route, exploiting portal venous
106 drainage for enhanced hepatic bioavailability of active constituents.

107 **Exercise and Yoga:** Structured daily sessions targeting large muscle groups to augment
108 peripheral insulin sensitivity through non-insulin-dependent GLUT-4 translocation. Pranayama
109 practices were incorporated to support autonomic regulation.

110 Antidiabetic medications were tapered systematically based on glycemic response during the
111 program, under medical supervision.

112 **2.4 Data Collection and Outcome Measures**

113 Baseline (pre-intervention) and post-intervention measurements were recorded at program
114 completion. Primary outcome: HbA1c (%). Secondary outcomes: fasting blood glucose (RBS,
115 mg/dL), body weight (kg), body mass index (kg/m²), abdominal girth (cm), systolic blood
116 pressure (SBP, mmHg), diastolic blood pressure (DBP, mmHg), resting heart rate (HR, bpm),
117 and antidiabetic medication status. Medication reduction was classified as: complete
118 discontinuation (Reduction Percentage = 1), partial reduction (0.3–0.5), or no change.
119 Panchakarma dose intensity was quantified as DonePK — the actual number of Panchakarma
120 sessions completed.

121 **2.5 Statistical Analysis**

122 Data were analyzed using Python 3 (scipy.stats). Paired two-tailed t-tests assessed pre-post
123 differences for all continuous outcomes. Results are reported as mean \pm standard deviation with
124 95% confidence intervals for mean change and corresponding p-values. Pearson correlation
125 coefficients evaluated the association between Panchakarma dose intensity (DonePK) and
126 change in HbA1c and body weight. Statistical significance was set at $p < 0.05$. Analyses were
127 restricted to participants with complete data for each outcome variable.

128

129 **3. RESULTS**

130 3.1 Participant Characteristics

131 Thirty-two patients met inclusion criteria (23 male, 9 female; mean age 46.1 ± 13.3 years).
132 Comorbidities included concomitant hypertension in 6 patients (18.8%), obesity in 11 (34.4%),
133 and coronary artery disease in 1 (3.1%). Mean Panchakarma sessions completed (DonePK):
134 10.3 ± 3.6 (range 5–19). Twenty-eight of 32 patients (87.5%) were on antidiabetic medications
135 at baseline. Table 1 presents participant characteristics.

136 **Table 1. Baseline Participant Characteristics (n = 32)**

Characteristic	Value
Age (years), mean \pm SD	46.1 ± 13.3
Sex, Male / Female	23 / 9
Concomitant hypertension, n (%)	6 (18.8%)
Concomitant obesity, n (%)	11 (34.4%)
On antidiabetic medications at baseline, n (%)	28 (87.5%)
DonePK (Panchakarma sessions), mean \pm SD	10.3 ± 3.6
DonePK range	5 – 19

137 3.2 Glycemic Outcomes

138 HbA1c declined significantly from $8.67 \pm 2.28\%$ to $7.47 \pm 1.68\%$ (mean reduction 1.19%, 95%
139 CI 0.67–1.71%, $p < 0.001$). Fifty percent of participants ($n = 16$) achieved HbA1c $< 7.0\%$ at
140 program completion, and 37.5% ($n = 12$) achieved HbA1c $< 6.5\%$. Clinically meaningful
141 glycemic response ($\geq 1\%$ absolute HbA1c reduction) was observed in 43.8% of participants ($n =$
142 14), and $\geq 2\%$ reduction in 21.9% ($n = 7$). Fasting blood glucose declined from 220.8 ± 106.3
143 mg/dL to 160.9 ± 64.3 mg/dL ($\Delta -59.9$ mg/dL, 95% CI 28.0–91.8, $p = 0.001$).

144 3.3 Anthropometric Outcomes

145 Body weight declined from 74.2 ± 16.4 kg to 70.0 ± 14.8 kg (mean reduction 4.28 kg, 95% CI
146 2.66–5.90, $p < 0.0001$). BMI fell from 27.6 ± 5.9 to 26.1 ± 5.2 kg/m² ($\Delta -1.52$ kg/m², $p < 0.0001$).
147 Abdominal girth ($n = 30$) decreased from 97.8 ± 20.5 cm to 93.0 ± 18.2 cm ($\Delta -4.73$ cm, 95% CI
148 2.96–6.51, $p < 0.0001$), indicating significant central adiposity reduction.

149 3.4 Hemodynamic and Autonomic Outcomes

150 Systolic blood pressure fell from 131.8 ± 14.5 to 124.7 ± 14.0 mmHg ($\Delta -7.16$ mmHg, 95% CI
151 1.97–12.35, $p = 0.011$) and diastolic blood pressure from 88.4 ± 12.4 to 82.0 ± 9.5 mmHg (Δ
152 -6.45 mmHg, 95% CI 2.14–10.77, $p = 0.006$). Resting heart rate decreased significantly from

153 88.4 ± 12.1 to 80.7 ± 12.2 bpm (Δ -7.68 bpm, 95% CI 4.24–11.11, $p < 0.001$). These findings
 154 suggest simultaneous improvement in hemodynamic load and autonomic tone.

155 3.5 Medication Reduction

156 Of the 28 patients on antidiabetic medications at baseline, 11 (39.3%) achieved complete
 157 medication discontinuation by program completion. A further 6 patients (21.4%) achieved partial
 158 dose reduction. Overall, 60.7% of medicated patients experienced measurable reduction in
 159 antidiabetic medication burden.

160 3.6 Dose-Intensity Relationship

161 Pearson correlation analysis revealed a significant inverse association between Panchakarma
 162 dose intensity (DonePK) and body weight reduction ($r = -0.46$, $p = 0.008$), indicating that
 163 participants completing more Panchakarma sessions achieved greater adiposity reduction. No
 164 significant correlation was observed between DonePK and HbA1c change ($r = -0.16$, $p =$
 165 0.376), suggesting that glycemic improvement was not linearly dose-dependent within the range
 166 of sessions delivered.

167 **Table 2. Pre- and Post-Intervention Cardiometabolic Outcomes (n = 32)**

Parameter	Baseline (Mean ± SD)	Post (Mean ± SD)	Δ Mean (95% CI)	p-value
HbA1c (%)	8.67 ± 2.28	7.47 ± 1.68	-1.19 (0.67–1.71)	< 0.001
Fasting Glucose (mg/dL)	220.8 ± 106.3	160.9 ± 64.3	-59.9 (28.0–91.8)	0.001
Weight (kg)	74.2 ± 16.4	70.0 ± 14.8	-4.28 (2.66–5.90)	< 0.0001
BMI (kg/m ²)	27.6 ± 5.9	26.1 ± 5.2	-1.52 (0.94–2.10)	< 0.0001
Abdominal Girth (cm)	97.8 ± 20.5	93.0 ± 18.2	-4.73 (2.96–6.51)	< 0.0001
SBP (mmHg)	131.8 ± 14.5	124.7 ± 14.0	-7.16 (1.97–12.35)	0.011
DBP (mmHg)	88.4 ± 12.4	82.0 ± 9.5	-6.45 (2.14–10.77)	0.006
Resting HR (bpm)	88.4 ± 12.1	80.7 ± 12.2	-7.68 (4.24–11.11)	< 0.001

168 *SBP = systolic blood pressure; DBP = diastolic blood pressure; HR = heart rate; CI = confidence interval*

169

170 4. DISCUSSION

171 The principal finding of this study is that the CDC program produced simultaneous, statistically
172 significant improvements across all measured cardiometabolic domains — glycaemia, adiposity,
173 blood pressure, and resting heart rate — in a semi-rural T2DM cohort. This breadth of effect
174 distinguishes multimodal Ayurvedic intervention from pharmacological monotherapies, which
175 characteristically address individual risk factors rather than the integrated pathophysiology of
176 cardiometabolic disease.

177 The mean HbA1c reduction of 1.19% (95% CI 0.67–1.71%) is comparable to that reported with
178 GLP-1 receptor agonists and SGLT-2 inhibitors in real-world studies, and exceeds the clinically
179 meaningful threshold of 0.5% reduction established by major diabetology guidelines.^{10,11} Fifty
180 percent of participants achieved an HbA1c < 7.0% — the standard American Diabetes
181 Association target — and 37.5% achieved < 6.5%. These proportions are notable given the
182 program duration and the mean baseline HbA1c of 8.67%, reflecting a predominantly poorly-
183 controlled cohort at enrolment.

184 The significant reductions in systolic (7.16 mmHg) and diastolic blood pressure (6.45 mmHg)
185 are clinically relevant: meta-analyses of antihypertensive trials indicate that a 5 mmHg SBP
186 reduction reduces the risk of major cardiovascular events by approximately 10%.¹² The
187 mechanisms underlying blood pressure reduction in the CDC program are likely multifactorial,
188 including weight loss (mean 4.28 kg), reduced sympathetic tone via yoga and pranayama, direct
189 vasodilatory effects of DashmulaKwath Swedan, and the natriuretic properties of glycyrrhizin at
190 the doses delivered via Basti. Although glycyrrhizin can elevate blood pressure at high systemic
191 exposures via mineralocorticoid pathways, the portal-hepatic delivery route via Basti involves
192 substantially lower systemic bioavailability, and blood pressure reduction was consistently
193 observed in this cohort.

194 The reduction in resting heart rate by 7.68 bpm ($p < 0.001$) is a finding of particular clinical
195 significance. Elevated resting heart rate (>80 bpm) is an independent predictor of
196 cardiovascular mortality in T2DM and the general population, with each 10 bpm increment
197 associated with an approximately 9% increase in all-cause mortality risk.^{13,14} The mean baseline
198 heart rate in our cohort was 88.4 bpm — above the prognostically adverse threshold. Post-
199 intervention, the mean fell to 80.7 bpm. This improvement likely reflects enhanced
200 parasympathetic tone through pranayama and yoga, improved cardiovascular fitness, reduced
201 sympathetic activation secondary to weight loss, and reduced insulin resistance (which itself
202 elevates sympathetic activity via central mechanisms).

203
204 The dose-response relationship between Panchakarma sessions and weight reduction ($r =$
205 -0.46 , $p = 0.008$) provides mechanistic support for the thermogenic and lipolytic contribution of
206 the Panchakarma procedures themselves, independent of dietary and exercise components.
207 This is consistent with prior findings from CDC program cohorts in urban settings.⁹ The absence
208 of a significant dose-response for HbA1c may reflect a threshold effect: glycemic improvement
209 occurs with even low-intensity intervention (dietary restriction alone is powerfully hypoglycemic
210 at 800 kcal), whereas progressive adiposity reduction — which involves deeper restructuring of
211 adipose tissue metabolism — benefits more continuously from increasing procedural intensity.

212 The medication outcomes in this cohort are notable: 39.3% of medicated patients achieved
213 complete antidiabetic medication discontinuation, and an additional 21.4% achieved dose
214 reduction. This deprescribing outcome has relevance not only for patient quality of life but for
215 the economic burden of chronic pharmacotherapy, which is a significant contributor to out-of-
216 pocket healthcare expenditure in rural Maharashtra.

217 Limitations of this study include the pre-post, single-arm design without a randomized control
218 group, the modest sample size ($n = 32$), and the semi-rural clinic setting that may limit
219 generalizability. The absence of lipid profile data in this dataset precluded evaluation of
220 dyslipidemia outcomes. Longer-term follow-up data would be required to evaluate durability of
221 benefits. Future studies should incorporate randomized designs, control groups, and extended
222 follow-up periods to establish causal attribution and duration of effect.

223 **5. CONCLUSION**

224 The CDC program produced clinically meaningful and statistically significant improvements
225 across glycemic, anthropometric, hemodynamic, and autonomic parameters in a semi-rural
226 T2DM cohort in the Marathwada region of Maharashtra, India. The simultaneous reduction in
227 HbA1c, body weight, abdominal girth, systolic and diastolic blood pressure, and resting heart
228 rate — together with a high rate of antidiabetic medication discontinuation — positions the CDC
229 program as a comprehensive cardiometabolic risk reduction strategy rather than a disease-
230 specific glycemic intervention. The dose-dependent relationship between Panchakarma
231 sessions and adiposity reduction provides mechanistic support for the procedural component's
232 contribution to metabolic outcomes. These findings add to a growing evidence base supporting
233 the global applicability of structured Ayurvedic multimodal interventions in the management of
234 T2DM and associated cardiovascular risk.

235 **REFERENCES**

- 236 1. Sun H, Saeedi P, Karuranga S, et al. IDF Diabetes Atlas: Global, regional and country-level diabetes
237 prevalence estimates for 2021 and projections for 2045. *Diabetes Res Clin Pract.* 2022;183:109119.
- 238 2. Mohan V, Sandeep S, Deepa R, Shah B, Varghese C. Epidemiology of type 2 diabetes: Indian
239 scenario. *Indian J Med Res.* 2007;125(3):217-230.
- 240 3. American Diabetes Association. 10. Cardiovascular disease and risk management. *Diabetes Care.*
241 2023;46(Suppl 1):S158-S190.
- 242 4. Holman RR, Paul SK, Bethel MA, Matthews DR, Neil HA. 10-year follow-up of intensive glucose control
243 in type 2 diabetes. *N Engl J Med.* 2008;359(15):1577-1589.
- 244 5. Maruthur NM, Tseng E, Hutfless S, et al. Diabetes medications as monotherapy or metformin-based
245 combination therapy for type 2 diabetes. *Ann Intern Med.* 2016;164(11):740-751.
- 246 6. Agnivesha. *Charaka Samhita. Chikitsasthana, Chapter 6 (Prameha Chikitsa).* Varanasi: Chaukhamba
247 Sanskrit Pratishthan; 2004.
- 248 7. Sushruta. *Sushruta Samhita. Nidanasthana, Chapter 6.* Varanasi: ChaukhambaVisvabharati; 2008.
- 249 8. Gond B. Glycemic and anthropometric outcomes of the Comprehensive Diabetic Care program:
250 evidence from an urban cohort in Maharashtra. [Institutional Research Report, Madhavbaug Clinics,
251 2023].
- 252 9. Gond B. Panchakarma dose-intensity and adiposity reduction in type 2 diabetes: findings from the CDC
253 program. [Institutional Research Report, Madhavbaug Clinics, 2023].
- 254 10. Zinman B, Wanner C, Lachin JM, et al. Empagliflozin, cardiovascular outcomes, and mortality in type
255 2 diabetes. *N Engl J Med.* 2015;373(22):2117-2128.
- 256 11. Marso SP, Daniels GH, Brown-Frandsen K, et al. Liraglutide and cardiovascular outcomes in type 2
257 diabetes. *N Engl J Med.* 2016;375(4):311-322.
- 258 12. Ettehad D, Emdin CA, Kiran A, et al. Blood pressure lowering for prevention of cardiovascular disease
259 and death: a systematic review and meta-analysis. *Lancet.* 2016;387(10022):957-967.
- 260 13. Fox K, Borer JS, Camm AJ, et al. Resting heart rate in cardiovascular disease. *J Am Coll Cardiol.*
261 2007;50(9):823-830.
- 262 14. Cooney MT, Vartiainen E, Laatikainen T, et al. Elevated resting heart rate is an independent risk
263 factor for cardiovascular disease in healthy men and women. *Am Heart J.* 2010;159(4):612-619.