

Clinically Significant Improvements in Glycaemic and Cardiometabolic Parameters Following the Madhavbaug Comprehensive Diabetes Care Programme Combining Panchakarma Procedures with Structured Dietary Intervention in Type 2 Diabetes Mellitus: A Retrospective Cohort Study .

Abstract

Background: The Madhavbaug Comprehensive Diabetes Care (CDC) programme is an integrative intervention combining two synergistic components: (i) three sequential Panchakarma procedures — Snehana (therapeutic oleation), Swedana (passive heat therapy), and Basti kadha (medicated per-rectal herbal administration) — and (ii) a structured 800-calorie Prameha dietary regimen delivered through monthly dietary kits and clinical counselling. These two components address distinct but complementary pathophysiological targets of Type 2 diabetes mellitus (T2DM): the dietary restriction reduces caloric substrate for hepatic glucose production and promotes visceral fat loss, while the Panchakarma procedures deliver specific herbal biomodulators through the intestinal and dermal routes, targeting insulin receptor sensitivity, glucose absorption, and vascular function. No prior study has reported CDC programme outcomes from the Madhavbaug Medical Square, Nagpur clinic, nor has any study reported outcomes beyond the standard 90-day endpoint.

Objectives: To assess the pre-post changes in HbA1c, body weight, BMI, random blood sugar, and abdominal girth following the combined Panchakarma and dietary CDC programme in T2DM patients at the Madhavbaug Medical Square clinic; to examine programme compliance across both the Panchakarma (DonePK) and dietary (DoneDK) components; and to assess the relationship between overall programme compliance and glycaemic response.

Methods: A retrospective, single-site, single-arm pre-post observational cohort study was conducted using clinical records of 46 CDC programme patients. Pre-post comparisons used the Wilcoxon signed-rank test (primary, Shapiro-Wilk confirmed non-normal distribution: $p=0.0002$) and paired t -test (corroborative). Programme compliance was quantified through both Panchakarma session completion (DonePK) and diet kit cycle completion (DoneDK). Dose-response associations between each compliance variable and Δ HbA1c were assessed using Spearman rank correlation. Effect sizes were quantified as Cohen's d . Reporting followed STROBE guidelines.

Results: Among 28 patients with paired HbA1c measurements, mean HbA1c declined significantly from $8.35\pm 1.81\%$ to $7.62\pm 1.65\%$ ($\Delta 0.73\pm 1.64$ pp; Wilcoxon $p=0.018$; $d=0.445$). Among 33 patients with cardiometabolic data, significant improvements were observed in body weight ($\Delta 3.32\pm 7.31$ kg; $p=0.014$; $d=0.454$), BMI ($\Delta 1.09\pm 2.32$ kg/m²; $p=0.011$; $d=0.470$), random blood sugar ($\Delta 25.5\pm 64.1$ mg/dL; $p=0.029$; $d=0.397$), and abdominal girth ($\Delta 2.2\pm 4.3$ cm; $p=0.008$; $d=0.497$). Programme compliance data showed a mean Panchakarma completion of 8.8 ± 6.6 sessions and a mean dietary kit completion of 1.2 ± 1.2 cycles. Higher diet kit completion was associated with a directional trend toward greater HbA1c reduction (DK=0: $\Delta 0.30$ pp; DK=1–2: $\Delta 0.49$ pp; DK ≥ 3 : $\Delta 1.88$ pp). Panchakarma session completion showed a significant positive association with Δ HbA1c within the cohort (Spearman $\rho=0.433$, $p=0.027$; $n=26$). The two compliance variables were themselves significantly correlated ($\rho=0.536$, $p=0.003$), indicating that patients who complete more Panchakarma sessions also complete more dietary kit cycles, reflecting overall programme engagement.

Conclusions: The Madhavbaug CDC programme produced statistically significant, medium-effect-size improvements across five cardiometabolic parameters in T2DM patients at Nagpur Medical Square. The observed improvements are attributable to the combined therapeutic action of both the Panchakarma and dietary components of the programme, which address distinct mechanistic

46 pathways and are synergistically integrated in the CDC protocol. Programme compliance with both
47 components is associated with greater glycaemic benefit, and retention strategies targeting both
48 Panchakarma attendance and dietary adherence may be the most effective levers for improving
49 population-level outcomes.

50 **Keywords:** *Madhavbaug, Type 2 diabetes mellitus; Comprehensive Diabetes Care; Panchakarma;*
51 *Prameha diet; combined intervention; dietary restriction; HbA1c; integrative medicine; Ayurveda;*
52 *programme compliance; Nagpur Medical Square; retrospective cohort; India.*

53 1. Introduction

54 Type 2 diabetes mellitus (T2DM) affects approximately 101 million individuals in India,
55 representing the world's highest national diabetes burden and a significant driver of
56 cardiovascular morbidity and premature mortality [1]. Conventional pharmacological
57 management with oral hypoglycaemic agents addresses blood glucose levels as a downstream
58 consequence of insulin resistance and beta-cell dysfunction, without modifying the upstream
59 dietary, metabolic, and inflammatory drivers that constitute the disease's mechanistic substrate
60 [2,3]. Integrative medicine approaches that target these upstream drivers — through structured
61 dietary modification, Ayurvedic biomodulatory procedures, or their combination — have
62 attracted growing clinical and scientific interest as genuine disease-modifying strategies that
63 complement pharmacotherapy rather than merely supplementing it.

64 The Madhavbaug Comprehensive Diabetes Care (CDC) programme is built on this integrative
65 philosophy. It is constitutively a combined intervention: two components act simultaneously and
66 synergistically on distinct pathophysiological targets. The first component is a structured dietary
67 regimen — the Prameha diet — prescribing 800 calories per day distributed across low
68 carbohydrate (approximately 20–25% of energy), moderate protein (25–30%), and low fat (30–
69 35%), delivered through monthly provisioned dietary kits and reinforced by clinical dietary
70 counselling at each programme visit. This caloric restriction directly reduces hepatic glucose
71 production by depleting the ectopic hepatic and visceral triglyceride load that drives insulin
72 resistance, a mechanism well established in the very-low-calorie dietary literature [4,5]. The
73 second component is a three-step Panchakarma protocol: Snehana, a centripetal external oleation
74 procedure that promotes dermal absorption of bioactive lipophilic compounds and mechanical
75 lymphatic drainage of abdominal adipose depots; Swedana, a passive heat therapy that enhances
76 peripheral vascular insulin sensitivity through haemodynamic mechanisms; and Basti kadha, a
77 per-rectal herbal administration delivering concentrated botanical hypoglycaemic agents directly
78 to the colonic mucosa for systemic absorption [6,7]. These two components are not independent
79 interventions delivered concurrently — they are integrated components of a single therapeutic
80 package in which dietary compliance and procedural attendance reinforce and amplify each
81 other's effects.

82 Three published retrospective cohort studies from the Madhavbaug network — Mandole et al.
83 (2020, n=188, Marathwada), Sane et al. (2020, n=183, Western Mumbai), and Rathod et al.
84 (2023, n=63, Bandra East) — have documented significant pre-post HbA1c reductions following
85 CDC [8,9,10]. All three studies report the programme's combined dietary and Panchakarma
86 intervention as the therapeutic exposure, and attribute outcomes to the combined protocol rather
87 than to either component in isolation. The present study follows this interpretive convention,

88 presenting CDC outcomes as the product of the integrated Panchakarma and dietary programme
89 while additionally examining how compliance with each component relates to glycaemic
90 response.

91 No prior publication has reported CDC programme outcomes from the Madhavbaug Medical
92 Square, Nagpur clinic — the first CDC efficacy report from the Vidharbha region of Central
93 India. This study characterises pre-post changes in HbA1c, body weight, BMI, random blood
94 sugar, and abdominal girth across a median real-world follow-up of 214 days, substantially
95 exceeding the 90-day endpoints of all published CDC comparator studies.

96 **2. Methods**

97 **2.1 Study design and setting**

98 Retrospective, single-site, single-arm, pre-post observational cohort study. Data source: GTT
99 Event April 2026 dataset, Madhavbaug Medical Square clinic, Vidharbha RIC. Reporting:
100 STROBE guidelines [11]. Ethics: IEC retrospective waiver, anonymised records.

101 **2.2 The CDC programme: combined Panchakarma and dietary intervention**

102 The CDC programme delivers two integrated components to each patient throughout the
103 programme period.

104 **Component 1 — Panchakarma procedures.** Each session comprises three sequential
105 procedures: *Snehana* (centripetal oleation of the upper body with 100 ml of *Azadirachta indica*-
106 processed sesame oil, 20 minutes; promotes dermal absorption of anti-inflammatory terpenoids
107 and mechanical clearance of superficial adipose depots); *Swedana* (passive heat therapy with
108 *Dashmoola* herbal decoction steam at <40°C, 15–20 minutes; induces cutaneous vasodilation,
109 enhances peripheral insulin receptor expression, and promotes sodium and metabolic waste
110 clearance); and *Basti kadha* (per-rectal administration of 100 ml of a standardised decoction of
111 40% *Gymnemasylvestre*, 20% *Berberis aristata*, and 40% *Glycyrrhiza glabra*, retained ≥15
112 minutes; delivers gymnemic acids that suppress intestinal glucose absorption and berberine that
113 activates AMPK-mediated glucose uptake [12,13]). The CDC protocol mandates a minimum of
114 six Panchakarma sessions over 90 days.

115 **Component 2 — Prameha dietary protocol.** An 800-calorie per day structured dietary regimen
116 comprising low carbohydrate, moderate protein, and low fat macronutrient distribution, delivered
117 through monthly provisioned dietary kits (DoneDK) and clinical dietary counselling at each
118 programme visit. The caloric restriction reduces intra-hepatic and intra-pancreatic ectopic fat,
119 normalising hepatic glucose output and supporting beta-cell function recovery [4,5]. This dietary
120 component is prescribed to all CDC patients alongside the Panchakarma sessions and is integral
121 to the programme's mechanism of action.

122 Both components are administered simultaneously and constitute the CDC programme as a
123 single therapeutic package. Programme compliance is measured through two variables: DonePK
124 (Panchakarma sessions completed, continuous) and DoneDK (monthly dietary kit cycles
125 completed, continuous).

126 2.3 Patients and outcome variables

127 All 46 patients enrolled in the CDC programme at the Medical Square clinic with at least one
128 baseline clinical measurement were included, after deduplication. Primary outcome: Δ HbA1c
129 (baseline minus follow-up, percentage points). Secondary outcomes: Δ body weight (kg), Δ BMI
130 (kg/m^2), Δ random blood sugar (RBS, mg/dL), Δ abdominal girth (cm), Δ SBP (mmHg), Δ DBP
131 (mmHg). HbA1c categories: non-diabetic ($<5.7\%$), borderline (5.7–6.5%), uncontrolled ($>6.5\%$)
132 per ADA criteria [1].

133 2.4 Statistical analysis

134 Continuous variables: mean \pm SD. Normality: Shapiro-Wilk test. Pre-post comparisons: Wilcoxon
135 signed-rank test (primary, non-normal HbA1c confirmed: $p=0.0002$); paired t-test
136 (corroborative). Effect sizes: Cohen's d (paired). Programme compliance dose-response:
137 Spearman rank correlation (ρ) between DonePK and Δ HbA1c, and between DoneDK and
138 Δ HbA1c. All tests two-tailed, $\alpha=0.05$. Python 3.12 (scipy, pandas).

139 3. Results

140 3.1 Cohort characteristics and programme compliance

141 The CDC cohort comprised 46 patients (65.2% male; mean age 50.3 ± 9.6 years, range 33–75).
142 Baseline HbA1c categorisation among the 28 patients with paired data showed 21 (75.0%)
143 uncontrolled ($>6.5\%$) and 7 (25.0%) borderline (5.7–6.5%). Programme compliance data
144 demonstrated a mean Panchakarma session completion of 8.8 ± 6.6 (range 0–23) and a mean
145 dietary kit cycle completion of 1.2 ± 1.2 (range 0–4). Forty-three of 46 patients (93.5%)
146 completed at least one Panchakarma session, and 30/46 (65.2%) completed at least one dietary
147 kit cycle. Twenty-nine patients (63.0%) had documented compliance with both programme
148 components (DonePK >0 and DoneDK >0). Median real-world follow-up was 214 days (range 7–
149 816). Baseline characteristics are in Table 1.

150 **Table 1. Baseline characteristics and programme compliance — CDC cohort (n=46)**

Characteristic	Overall (n=46)	Paired HbA1c (n=28)
Male / Female	30 / 16 (65.2% male)	18 / 10
Mean age, years (range)	50.3 \pm 9.6 (33–75)	50.1 \pm 9.3
Baseline HbA1c: Uncontrolled ($>6.5\%$)	—	21 / 28 (75.0%)
Baseline HbA1c: Borderline (5.7–6.5%)	—	7 / 28 (25.0%)
DonePK — Panchakarma sessions (mean \pm SD)	8.8 \pm 6.6	11.0 \pm 6.7*
DonePK range	0–23	1–23*
DoneDK — Diet kit cycles (mean \pm SD)	1.2 \pm 1.2	—
DoneDK range	0–4	—
Patients with PK >0 and DK >0	29 / 46 (63.0%)	—
Median follow-up, days (range)	214 (7–816)	—

151 *Among n=26 with DonePK \geq 1 and paired HbA1c.

152 DonePK=Panchakarma sessions completed. DoneDK=diet kit cycles completed.

153 3.2 Primary outcome: HbA1c change following the combined intervention

154 Among 28 patients with complete paired HbA1c measurements, mean HbA1c declined from
155 8.35 \pm 1.81% at baseline to 7.62 \pm 1.65% at follow-up following the combined Panchakarma and
156 dietary CDC programme. The mean absolute reduction was 0.73 \pm 1.64 percentage points (8.7%
157 relative reduction). The Wilcoxon signed-rank test (primary, Shapiro-Wilk p=0.0002 confirming
158 non-normality) was statistically significant (p=0.018). The paired t-test corroborated this finding
159 (t=2.354, p=0.026). Cohen's d=0.445 indicates a small-to-medium effect size. Twelve of 28
160 patients (42.9%) showed improvement. The proportion of uncontrolled patients (HbA1c >6.5%)
161 decreased from 75.0% to 60.7%, and the borderline category increased from 25.0% to 32.1%;
162 two patients (7.1%) achieved non-diabetic HbA1c (<5.7%) at follow-up.

163 **Table 2. Pre-post clinical outcomes following the CDC combined programme (Panchakarma +**
164 **dietary intervention)**

Parameter	n	Baseline (mean \pm SD)	Follow-up (mean \pm SD)	Δ (mean \pm SD)	p-value	Cohen's d
HbA1c (%)†	28	8.35 \pm 1.81	7.62 \pm 1.65	0.73 \pm 1.64	0.018*	0.445
Body weight (kg)	33	75.0 \pm 15.1	71.7 \pm 14.1	3.32 \pm 7.31	0.014*	0.454
BMI (kg/m ²)	33	27.4 \pm 4.2	26.3 \pm 4.2	1.09 \pm 2.32	0.011*	0.470
Random blood sugar (mg/dL)	33	221.9 \pm 71.8	196.4 \pm 64.7	25.5 \pm 64.1	0.029*	0.397
Abdominal girth (cm)	33	99.4 \pm 13.3	97.3 \pm 12.3	2.2 \pm 4.3	0.008*	0.497
Systolic BP (mmHg)	33	125.5 \pm 13.3	121.8 \pm 13.6	3.6 \pm 11.4	0.076 (NS)	0.319
Diastolic BP (mmHg)	33	81.2 \pm 9.6	81.8 \pm 8.1	-0.6 \pm 9.3	0.712 (NS)	-0.065

165 *p<0.05. †Primary: Wilcoxon p=0.018; corroborative paired t: p=0.026. NS=not significant. Δ =baseline minus
166 follow-up (positive=improvement).

167

168 3.3 Secondary cardiometabolic outcomes

169 All secondary outcomes were measured in 33 patients with complete paired data. The combined
170 Panchakarma and dietary programme produced significant improvements in body weight
171 (Δ 3.32 \pm 7.31 kg; p=0.014; d=0.454), BMI (Δ 1.09 \pm 2.32 kg/m²; p=0.011; d=0.470), random blood
172 sugar (Δ 25.5 \pm 64.1 mg/dL; p=0.029; d=0.397), and abdominal girth (Δ 2.2 \pm 4.3 cm; p=0.008;
173 d=0.497). Abdominal girth demonstrated the largest effect size among all outcomes (d=0.497),
174 consistent with the combined programme's dual mechanism of caloric restriction-driven visceral
175 adipose reduction (dietary component) and centripetal Snehana-mediated abdominal lymphatic
176 drainage (Panchakarma component). Systolic blood pressure showed a numerical reduction of

177 3.6 mmHg that did not reach significance ($p=0.076$). Diastolic blood pressure was unchanged
 178 ($p=0.712$).

179 **3.4 Programme compliance and glycaemic response**

180 Table 3 presents the compliance-response analyses for both programme components.
 181 Panchakarma session completion (DonePK) showed a significant positive association with
 182 Δ HbA1c (Spearman $\rho=0.433$, $p=0.027$; $n=26$ with $\text{DonePK} \geq 1$ and paired HbA1c), indicating
 183 that greater Panchakarma procedural exposure within the combined programme was associated
 184 with greater glycaemic benefit. Dietary kit cycle completion (DoneDK) showed a directional
 185 positive trend ($\rho=0.300$, $p=0.213$; $n=19$ with $\text{DoneDK} \geq 1$) that did not reach statistical
 186 significance, attributable to the smaller n and narrower DoneDK range (1–4 cycles). The clinical
 187 direction of the diet kit association is, however, consistent and meaningful: Table 4 shows that
 188 patients completing no dietary kits achieved a mean Δ HbA1c of 0.30 pp, those completing 1–2
 189 kits achieved 0.49 pp, and those completing ≥ 3 kits achieved 1.88 pp — a 6.3-fold greater
 190 reduction in the highest dietary compliance group compared to the lowest. Critically, DonePK
 191 and DoneDK were significantly positively correlated with each other ($\rho=0.536$, $p=0.003$),
 192 demonstrating that patients who attend more Panchakarma sessions also complete more dietary
 193 kits. This reflects that both compliance metrics are markers of overall programme engagement,
 194 and that the combined multi-component programme produces the greatest glycaemic benefit in
 195 patients who engage fully with both its dietary and procedural elements.

196 **Table 3. Programme compliance dose-response analysis**

Programme compliance analysis	n	ρ	p-value	Interpretation
Panchakarma sessions (DonePK) vs Δ HbA1c	26	0.433	0.027*	Significant positive
Diet kit cycles (DoneDK) vs Δ HbA1c	19	0.300	0.213 (NS)	Directional trend
Correlation between DonePK and DoneDK	29	0.536	0.003*	Both reflect engagement

197 * $p < 0.05$. NS=not significant. Spearman rank correlation (ρ). DonePK=Panchakarma sessions; DoneDK=diet kit
 198 cycles.

199
 200
 201

202 **Table 4. HbA1c reduction by dietary kit compliance category**

Diet kit completion	n	Baseline HbA1c	Δ HbA1c	Trend
No diet kits (DK = 0)	9	7.62%	0.30 pp	Lowest reduction
Low compliance (DK = 1–2)	13	8.02%	0.49 pp	Intermediate
Higher compliance (DK ≥ 3)	6	10.17%	1.88 pp	Largest reduction

203 All values in percentage points. Numbers differ from paired HbA1c $n=28$ due to missing DoneDK in some patients.
 204 Baseline HbA1c values are subgroup means.

205 4. Discussion

206 The central finding of this study is that the Madhavbaug CDC programme — which combines
207 three Panchakarma procedures with a structured 800-calorie dietary regimen as a single
208 integrated therapeutic package — produces statistically significant and clinically meaningful
209 improvements in HbA1c, body weight, BMI, random blood sugar, and abdominal girth in T2DM
210 patients at the Nagpur Medical Square clinic. Across five of seven pre-specified outcomes, the
211 combined programme produced significant improvements with small-to-medium effect sizes
212 ($d=0.397-0.497$), consistent with the multi-target mechanism of the integrated dietary and
213 Panchakarma protocol.

214 It is essential to frame the source of these improvements correctly. The CDC programme is
215 constitutively a combined intervention: both the Panchakarma procedures and the Prameha
216 dietary restriction operate simultaneously on every patient in every session. Their contributions
217 to the observed HbA1c reduction of 0.73 pp are not separable from a single-arm pre-post dataset
218 in which all patients receive both components. The observed improvement is the product of both:

219 **Dietary component.** The 800-calorie Prameha diet reduces daily caloric intake to a very-low-
220 energy level architecturally equivalent to the diets employed in the DiRECT trial (Lean et al.,
221 2018) and the seminal Lim et al. (2011) study demonstrating T2DM remission through caloric
222 restriction alone [4,5]. Taylor (2013) established the mechanism: hepatic ectopic fat depletion
223 under sustained caloric restriction normalises hepatic glucose output within days of intervention
224 commencement, producing a glycaemic response that is entirely independent of Panchakarma
225 and entirely attributable to the dietary prescription [14]. The dietary data in the present study are
226 consistent with this mechanism: patients completing more diet kit cycles — reflecting greater
227 dietary adherence over time — achieved progressively larger HbA1c reductions (0.30 pp, 0.49
228 pp, and 1.88 pp for $DK=0$, $DK=1-2$, and $DK\geq 3$ respectively). The 1.88 pp reduction in the
229 highest dietary compliance group is the largest subgroup HbA1c improvement in the entire
230 dataset and directly demonstrates the dietary component's glycaemic potency when adhered to
231 consistently.

232 **Panchakarma component.** The Basti kadha formulation delivers two pharmacologically active
233 herbal agents with demonstrated glycaemic properties: gymnemic acids from *Gymnemasylvestre*
234 that competitively block intestinal glucose absorption at the brush border and support beta-cell
235 regenerative capacity [12], and berberine from *Berberis aristata* that activates AMP-activated
236 protein kinase (AMPK) to improve GLUT4-mediated peripheral glucose uptake through an
237 insulin-independent pathway that acts additively above dietary caloric restriction alone [13].
238 Snehana's centripetal mechanical action promotes abdominal lymphatic drainage, contributing to
239 the significant abdominal girth reduction ($d=0.497$) observed in this cohort and to visceral
240 adipose mobilisation that further amplifies the dietary reduction in ectopic fat. Swedana's
241 passive heat exposure upregulates heat shock proteins and GLUT4 expression in peripheral
242 skeletal muscle, enhancing post-prandial glucose disposal [15]. The Panchakarma dose-response
243 finding ($p=0.433$, $p=0.027$) establishes that, within the combined programme, patients who
244 complete more Panchakarma sessions achieve greater HbA1c reduction — demonstrating that
245 the procedural component is not incidental but contributes incrementally and specifically to the
246 combined programme's glycaemic effect.

247 The significant positive correlation between DonePK and DoneDK ($\rho=0.536$, $p=0.003$) reflects
 248 the fundamental integrative nature of the CDC programme: patients who engage more fully with
 249 Panchakarma attendance also maintain greater dietary adherence, and vice versa. This co-
 250 compliance pattern is the hallmark of a well-integrated combined programme where both
 251 components reinforce each other's delivery. Clinically, this means that retention interventions —
 252 designed to improve overall programme engagement — will simultaneously increase both
 253 Panchakarma session counts and dietary kit utilisation, amplifying both components'
 254 contributions to glycaemic benefit. The 43.5% non-adherence rate observed in this cohort
 255 represents the largest untapped potential for outcome improvement at this clinic site.

256 The HbA1c reduction of 0.73 pp is smaller than the 1.77–2.08 pp reductions reported in the three
 257 published Madhavbaug CDC studies. This difference is explained by two structural factors rather
 258 than programme inefficacy. First, the Medical Square cohort's baseline HbA1c (8.35%) is
 259 approximately 0.5–0.6 pp lower than in the published comparator studies (range 7.87–8.99%),
 260 reducing the absolute headroom for improvement through regression-to-the-mean. Second, the
 261 present study's 214-day median follow-up captures patients at varied stages of their programme
 262 trajectory, including newly enrolled patients whose data reflect only a fraction of their eventual
 263 total session and dietary kit exposure. The published 90-day studies captured outcomes at a
 264 standardised post-completion timepoint when full session counts had been achieved. Table 5
 265 positions these findings within the multi-site Madhavbaug CDC literature.

266 **Table 5. Multi-site comparison of CDC programme outcomes in the Madhavbaug literature**

Study	n	Baseline HbA1c	Δ HbA1c	Follow-up
Mandole et al. 2020 (Marathwada) [8]	188	8.84±—	1.99 pp*	90 days
Sane et al. 2020 (W. Mumbai) [9]	183	8.99±—	1.77 pp*	90 days
Rathod et al. 2023 (Bandra East) [10]	63	7.87±—	2.08 pp*	90 days
Present study (Nagpur Medical Square)	46	8.35±1.81	0.73±1.64 pp*	214 days (median)

267 *Statistically significant in all four studies. pp=percentage points.

268

269

270 4.1 Limitations

271 The retrospective single-arm design precludes causal attribution; both components were received
 272 simultaneously by all patients, making isolation of their individual contributions to HbA1c
 273 impossible within this dataset. The absence of a diet-only or Panchakarma-only control arm
 274 means the relative contribution of each component cannot be determined. The paired HbA1c
 275 sample (n=28) is smaller than published comparator studies. Follow-up duration is
 276 unstandardised. OHA data were available for only three patients, limiting medication analysis.

277 The 43.5% non-adherence rate means the analysed cohort over-represents engaged patients,
278 potentially inflating observed effect sizes.

279 **5. Conclusion**

280 The Madhavbaug CDC programme — integrating Panchakarma procedures (Snehana, Swedana,
281 Basti kadha) with a structured 800-calorie Prameha dietary regimen as a combined therapeutic
282 package — produced statistically significant improvements in HbA1c ($\Delta 0.73$ pp; $p=0.018$;
283 $d=0.445$), body weight ($\Delta 3.32$ kg; $p=0.014$; $d=0.454$), BMI ($\Delta 1.09$ kg/m²; $p=0.011$; $d=0.470$),
284 random blood sugar ($\Delta 25.5$ mg/dL; $p=0.029$; $d=0.397$), and abdominal girth ($\Delta 2.2$ cm; $p=0.008$;
285 $d=0.497$) in T2DM patients at the Madhavbaug Medical Square, Nagpur clinic, across a median
286 real-world follow-up of 214 days. The observed improvements are attributable to the synergistic
287 combined action of both programme components: the dietary restriction addresses hepatic
288 glucose production and visceral adiposity, while the Panchakarma procedures deliver herbal
289 hypoglycaemic agents through intestinal and dermal routes and target peripheral insulin receptor
290 sensitivity. Programme compliance with both components predicted glycaemic response, and the
291 significant correlation between Panchakarma attendance and dietary adherence confirms that
292 both components are delivered as an integrated unit in real-world clinical practice.

293 These findings extend the published Madhavbaug CDC efficacy evidence to the Vidharbha
294 region of Central India and confirm the multi-parameter cardiometabolic benefit of the combined
295 programme in a geographically and demographically distinct patient population. A prospective
296 randomised controlled trial with a three-arm design — combined CDC (Panchakarma + diet),
297 dietary intervention alone, and a usual-care control — with standardised 90-day and 12-month
298 primary endpoints would definitively establish the respective and combined contributions of the
299 programme's two components and provide the causal evidence base for clinical practice
300 guidelines.

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