

Testosterone Replacement in Hypogonadal Men With Type 2 Diabetes and/or Metabolic Syndrome (the TIMES2 Study)

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OBJECTIVE—This study evaluated the effects of testosterone replacement therapy (TRT) on insulin resistance, cardiovascular risk factors, and symptoms in hypogonadal men with type 2 diabetes and/or metabolic syndrome (MetS).

RESEARCH DESIGN AND METHODS—The efficacy, safety, and tolerability of a novel transdermal 2% testosterone gel was evaluated over 12 months in 220 hypogonadal men with type 2 diabetes and/or MetS in a multicenter, prospective, randomized, double-blind, placebo-controlled study. The primary outcome was mean change from baseline in homeostasis model assessment of insulin resistance (HOMA-IR). Secondary outcomes were measures of body composition, glycemic control, lipids, and sexual function. Efficacy results focused primarily on months 0–6 (phase 1; no changes in medication allowed). Medication changes were allowed in phase 2 (months 6–12).

RESULTS—TRT reduced HOMA-IR in the overall population by 15.2% at 6 months ($P = 0.018$) and 16.4% at 12 months ($P = 0.006$). In type 2 diabetic patients, glycemic control was significantly better in the TRT group than the placebo group at month 9 (HbA_{1c}: treatment difference, -0.446% ; $P = 0.035$). Improvements in total and LDL cholesterol, lipoprotein a (Lpa), body composition, libido, and sexual function occurred in selected patient groups. There were no significant differences between groups in the frequencies of adverse events (AEs) or serious AEs. The majority of AEs ($>95\%$) were mild or moderate.

CONCLUSIONS—Over a 6-month period, transdermal TRT was associated with beneficial effects on insulin resistance, total and LDL-cholesterol, Lpa, and sexual health in hypogonadal men with type 2 diabetes and/or MetS.

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Low serum testosterone is common in men with type 2 diabetes and/or metabolic syndrome (MetS) and numerous studies have reported an association between testosterone deficiency and visceral obesity, insulin resistance (IR) and dyslipidemia (1–4). In men with type 2 diabetes, low testosterone is associated with a high prevalence of symptomatic hypogonadism (3), frequently due to hypogonadotrophic hypogonadism (5).

Small studies have demonstrated that testosterone replacement therapy (TRT) in hypogonadal men with and without type 2 diabetes is associated with reductions in IR, waist circumference, cholesterol, glycated hemoglobin (HbA_{1c}), and fasting plasma glucose (FPG) concentrations (6–9). Conversely, withdrawal of TRT in hypogonadal men leads to decreased insulin sensitivity (10). Furthermore, androgen suppression therapy for prostate cancer can result in alterations of individual cardiovascular risk factors and increases the occurrence of incident diabetes, myocardial infarction and sudden cardiac death (11,12). Epidemiologic studies have reported that low testosterone in men is associated with increased all-cause and cardiovascular mortality (13).

The TIMES2 (Testosterone replacement In hypogonadal men with either METabolic Syndrome or type 2 diabetes) study investigated the effects of transdermal TRT on IR, selected cardiovascular risk factors, and symptoms in hypogonadal men with MetS and/or type 2 diabetes. The safety and tolerability of a novel, metered-dose, transdermal 2% testosterone gel were also examined.

RESEARCH DESIGN AND METHODS

This 12-month, prospective, randomized, double-blind, placebo-controlled, multicenter study was conducted at 36 outpatient centers in Belgium, France, Germany, Italy, the Netherlands, Spain, Sweden, and the U.K. between February 2006 and March

2007. The study was conducted in accordance with the revised guidelines of the World Medical Association Declaration of Helsinki, the International Conference on Harmonization Standard Operating Procedures, and local laws and regulatory requirements. The study protocol was approved by independent ethics committees for each center. Participants provided written informed consent.

Men aged ≥ 40 years were eligible to enter the study if they had confirmed hypogonadism (early morning [0800–1000 h] total testosterone [TT] ≤ 11 nmol/L or free testosterone ≤ 255 pmol/L on two occasions ≥ 1 week apart), with at least two symptoms of hypogonadism (14) and fulfilled criteria for type 2 diabetes (15) and/or MetS (16) (Supplementary Data). Erectile dysfunction was not an inclusion criterion.

Exclusion criteria included TRT within 6 months of randomization, hormone-modulating therapies or topical/systemic glucocorticoids within 3 months of randomization, or insulin therapy; a history of current prostate or breast cancer; abnormal digital rectal examination suggestive of prostate carcinoma; severe symptomatic benign prostatic hyperplasia or elevated age-specific prostate-specific antigen (PSA) (17). For PSA levels and other exclusion criteria see the Supplementary Data. Concomitant medications (oral antidiabetic drugs [OADs], lipid-lowering therapies, antihypertensives) were continued if doses had remained stable for ≥ 3 months before randomization. The first 6 months of the study, during which no therapy changes were permitted unless the investigator deemed it necessary for good clinical practice, was designated Phase 1. The second 6-month period, when changes in medication were allowed, was designated Phase 2.

Subjects were randomized (1:1) to receive either 3 g metered-dose 2% testosterone gel (60 mg testosterone, Tosttran [also known as Fortigel, Tostrex, Itnogen, Foresta; ProStrakan, Galashiels, Scotland, U.K.]) or placebo gel once daily and stratified to treatment arms according to the presence of MetS only and type 2 diabetes with or without MetS. All individuals involved in monitoring, data management, or other study aspects were blinded to treatment.

Treatment was applied daily (0700–1000 h) to clean, dry, intact skin on the thighs or abdomen. Blood samples were taken 2 h after application. TT was measured at 2, 4, and 12 weeks with dose

adjustments made as follows: TT > 52 nmol/L, TRT dose reduced to 40 mg/day; TT < 17 nmol/L, dose increased to 80 mg/day. Dummy dose changes were performed in the placebo group to maintain blinding.

All laboratory assessments (Supplementary Data) were performed centrally by Pivotal Laboratories Ltd., York, U.K. Measurements were taken at baseline, then every 3 months for HbA_{1c}, PSA, lipids (total cholesterol [TC], HDL cholesterol, LDL cholesterol, triglycerides, lipoprotein a [Lpa]), hematology, biochemistry, testosterone, FPG (no 9-month measurement) and insulin (mean of three samples at 5-min intervals, no 9-month measurement). Free testosterone was calculated from TT, albumin, and sex hormone binding globulin, using the Vermeulen equation (18). Waist circumference, BMI, body composition (TANITA BF-300 body fat analyzer; TANITA Corporation, Tokyo, Japan), Aging Males Symptoms (AMS) score (19), and International Index of Erectile Function (IIEF) (20) were assessed at baseline, 6, and 12 months. Adverse events (AEs) were elicited from the patient at each visit by a non-leading question.

Statistical analysis

The primary end point was the difference between treatment groups in homeostasis model assessment (HOMA)-IR from baseline to months 6 and 12. Secondary end points were changes from baseline in HbA_{1c}, fasting insulin, FPG, lipid parameters, body composition, BMI, waist circumference, AMS and IIEF scores, AEs, and other safety parameters. HOMA of β -cell function (HOMA-B) was determined post hoc.

The safety population comprised all randomized patients who received ≥ 1 dose of study medication. The primary analysis of all end points was based on the intention-to-treat (ITT) population (all randomized patients). Two other populations were also studied: the per-protocol (PP) population (randomized patients who received ≥ 1 dose of study medication with no protocol violations during the first 6 months of the study) and the modified PP (mPP) population. This latter population was established post hoc because the PP population was substantially reduced in size after exclusion of patients who took unauthorized concomitant medications from the ITT population (Supplementary Data). Patients' eligibility for inclusion in the mPP

population was assessed on a case-by-case basis for each end point. Inclusion was allowed if there were no changes in drugs during the first 6 months that affected the end point in question. These were OADs and antiobesity drugs for HOMA-IR ($n = 130$); OADs, antiobesity drugs, β -blockers, and diuretics for body composition ($n = 147$); lipid-lowering drugs, β -blockers, and α -blockers for lipid profile ($n = 139$); OADs, antiobesity drugs, β -blockers, and diuretics for HbA_{1c} and FPG ($n = 140$); phosphodiesterase-5 inhibitors and β -blockers for AMS ($n = 152$) and IIEF ($n = 154$); and antihypertensive drugs for blood pressure ($n = 140$).

The study was designed to have a power of 95% to detect a mean difference in HOMA-IR change from baseline between TRT and placebo of 0.82 (SD 1.7) at 6 months. After completion of an interim analysis in 63 patients, the study power was revised to 80%, giving a final target completer sample of 136 patients. All statistical analyses (Chiltern International Ltd; Quantics, Newtown St Boswells, U.K.) were conducted using SAS version 8.2 (SAS Institute, Cary, NC) or later. Descriptive statistics were calculated for continuous variables, with 95% CI, SD, and P values. For comparison of efficacy outcomes between treatment groups (treatment difference [TD]), an ANCOVA model was applied, with disease category as a covariate (MetS and type 2 diabetes; MetS only; type 2 diabetes only). $P < 0.05$ was considered statistically significant for all analyses. No adjustment was made for multiple testing as all secondary end points were regarded as exploratory. For the primary end point and HOMA-B, a log transformation was applied for the ANCOVA and a last observation carried forward approach was used. Unless otherwise stated, all efficacy data were analyzed in the ITT population, all safety data were analyzed in the safety population, and all results are quoted as mean \pm SD.

RESULTS—Of 518 patients screened, 220 were randomized (testosterone $n = 108$; placebo $n = 112$) and received at least one dose of study drug (ITT and safety populations). Baseline demographic and clinical characteristics were comparable between groups (Table 1). Phase 1 was completed by 157 patients (71%), and Phases 1 and 2 were completed by 118 (54%) (Supplementary Fig. A1). Reasons for early withdrawal included protocol violation ($n = 28$), withdrawal of consent

Table 1—Baseline characteristics (safety population)

	Testosterone	Placebo	Total
<i>n</i>	108	112	220
Age (years)	59.9 ± 9.1 (37–77)	59.9 ± 9.4 (39–88)	59.9 ± 9.3 (37–88)
Race			
Caucasian	105 (97)	110 (98)	215 (98)
Asian	3 (3)	0	3 (1)
Afro-Caribbean	0	1 (1)	1 (0.5)
Other	0	1 (1)	1 (0.5)
Total serum testosterone (nmol/L)	9.2 ± 2.6 (3.8–20.1)	9.5 ± 3.3 (2.2–20.6)	9.4 ± 3.0 (2.2–20.6)
Free testosterone (pmol)	198.0 ± 49.3 (42–319)	202.4 ± 62.1 (35–364)	200.3 ± 56.1 (35–364)
Type 2 diabetes	68 (6)	69 (62)	137 (62)
MetS	88 (82)	88 (79)	176 (80)
Insulin resistance (HOMA-IR index)	5.9 ± 3.8 (1–20)	4.9 ± 3.3 (1–21)	5.4 ± 3.6 (1–21)
Hematocrit (L/L)	0.43 ± 0.04 (0.34–0.54)	0.43 ± 0.04 (0.33–0.51)	0.43 ± 0.04 (0.33–0.54)
Hemoglobin (g/dL)	14.9 ± 1.5 (11.0–18.6)	14.9 ± 1.3 (11.4–18.0)	14.9 ± 1.4 (11.0–18.6)
PSA (μg/L)	1.6 ± 1.8 (0.21–13.60)	1.2 ± 1.2 (0.20–6.38)	1.4 ± 1.5 (0.20–13.60)
FSH* (IU/L)	12.3 ± 13.4 (1.2–63.6)	12.5 ± 15.7 (1.2–78.9)	12.4 ± 14.5 (1.2–78.9)
FSH*			
Low	0	0	0
Normal	77 (71)	79 (71)	156 (71)
High	31 (29)	33 (29)	64 (29)
LH* (IU/L)	5.7 ± 5.7 (0.9–34.3)	5.3 ± 5.4 (0.6–36.1)	5.5 ± 5.5 (0.6–36.1)
LH*			
Low	0	2 (2)	2 (1)
Normal	86 (80)	92 (82)	178 (81)
High	22 (20)	18 (16)	40 (18)

Data are mean ± SD (range) or *n* (%) unless otherwise indicated. FSH, follicle stimulating hormone; LH, luteinizing hormone. *Screening values (not measured at baseline). Normal ranges: 0.7 IU/L ≤ FSH ≤ 11.1 IU/L; 0.8 IU/L ≤ LH ≤ 7.6 IU/L.

(*n* = 27), AEs (*n* = 26) and loss to follow-up (*n* = 9). Protocol was violated by 125 patients, giving a PP population of 95. Hypogonadotropic or secondary hypogonadism, defined as low or normal luteinizing hormone, occurred in 82% of subjects whereas 18% demonstrated primary hypogonadism (Supplementary Table A1). The majority of subjects (80%) had MetS at baseline, and 62% had type 2 diabetes. MetS and type 2 diabetes were both present in 44 and 40% of the TRT and placebo groups, respectively. Baseline serum TT values were 9.2 ± 2.6 nmol/L (TRT) and 9.5 ± 3.3 nmol/L (placebo). The majority of subjects (62.6%) qualified for study entry, meeting both TT and free testosterone criteria. Of the subjects, 12.3 and 23.7% qualified on the basis of TT <11 nmol/L or free testosterone <255 pmol/L, respectively.

Efficacy: Phase 1

At the end of Phase 1 (6 months), serum TT values in the ITT population had increased by 19.0 ± 22.1 nmol/L with TRT versus 0.1 ± 2.9 nmol/L with placebo (*P* < 0.001).

TRT reduced HOMA-IR by 15.2% (95% CI 3–26; *P* = 0.018) compared with

placebo at 6 months (Fig. 1). Type 2 diabetic patients demonstrated mean reductions in HOMA-IR of 16.0% (0–29; *P* = 0.049) compared with placebo. The difference between treatments in the MetS patients approached significance (TD 13.3% [–1 to 26]; *P* = 0.069) (Fig. 1). Post hoc analyses of HOMA-IR in patients without type 2 diabetes only, showed no difference between treatment groups (Supplementary Table A2). There were no significant differences between treatments in the overall population or the disease subgroups in HbA_{1c} (Fig. 1), fasting serum insulin, FPG, or HOMA-B (Table 2).

TRT had a number of beneficial effects on lipid profile. In the MetS subgroup, TRT was associated with significantly greater reductions from baseline in plasma levels of Lpa (TD –0.31 μmol/L [95% CI –0.543 to –0.082]; *P* = 0.008), TC (TD –0.336 mmol/L [–0.558 to –0.114]; *P* = 0.003) and LDL cholesterol (TD –0.210 mmol/L [95–0.374 to –0.047]; *P* = 0.012) than placebo (Table 2). TRT also reduced Lpa in the overall ITT population (TD –0.235 μmol/L [–0.431 to –0.039]; *P* = 0.019). HDL cholesterol decreased from baseline significantly more with TRT than placebo

in all three analysis groups (total population [TD –0.049 mmol/L (–0.094 to –0.004); *P* = 0.032], MetS [TD –0.058 mmol/L (–0.105 to –0.011); *P* = 0.016], type 2 diabetes [TD –0.062 mmol/L (–0.123 to –0.002); *P* = 0.043]). There were no significant effects of TRT on triglycerides, abdominal obesity, percentage body fat, BMI, or waist circumference.

In the overall ITT population, TRT was associated with significantly greater increases from baseline in total IIEF score (TD 4.868 [95% CI 0.644–9.092]; *P* = 0.024) and in the sexual desire domain (TD 0.800 [0.271–1.329]; *P* = 0.003) (Table 2). Treatment had no significant effect on erectile function, orgasmic function, or overall sexual satisfaction, although these three measures did show a consistent trend for greater improvement with TRT versus placebo. Change in AMS score was similar in both treatment groups.

Data for secondary end points in the mPP and PP populations are shown in Supplementary Tables A3 and A4, respectively. The small size of the PP population negated statistical power and the difference between groups in HOMA-IR reduction was not statistically significant

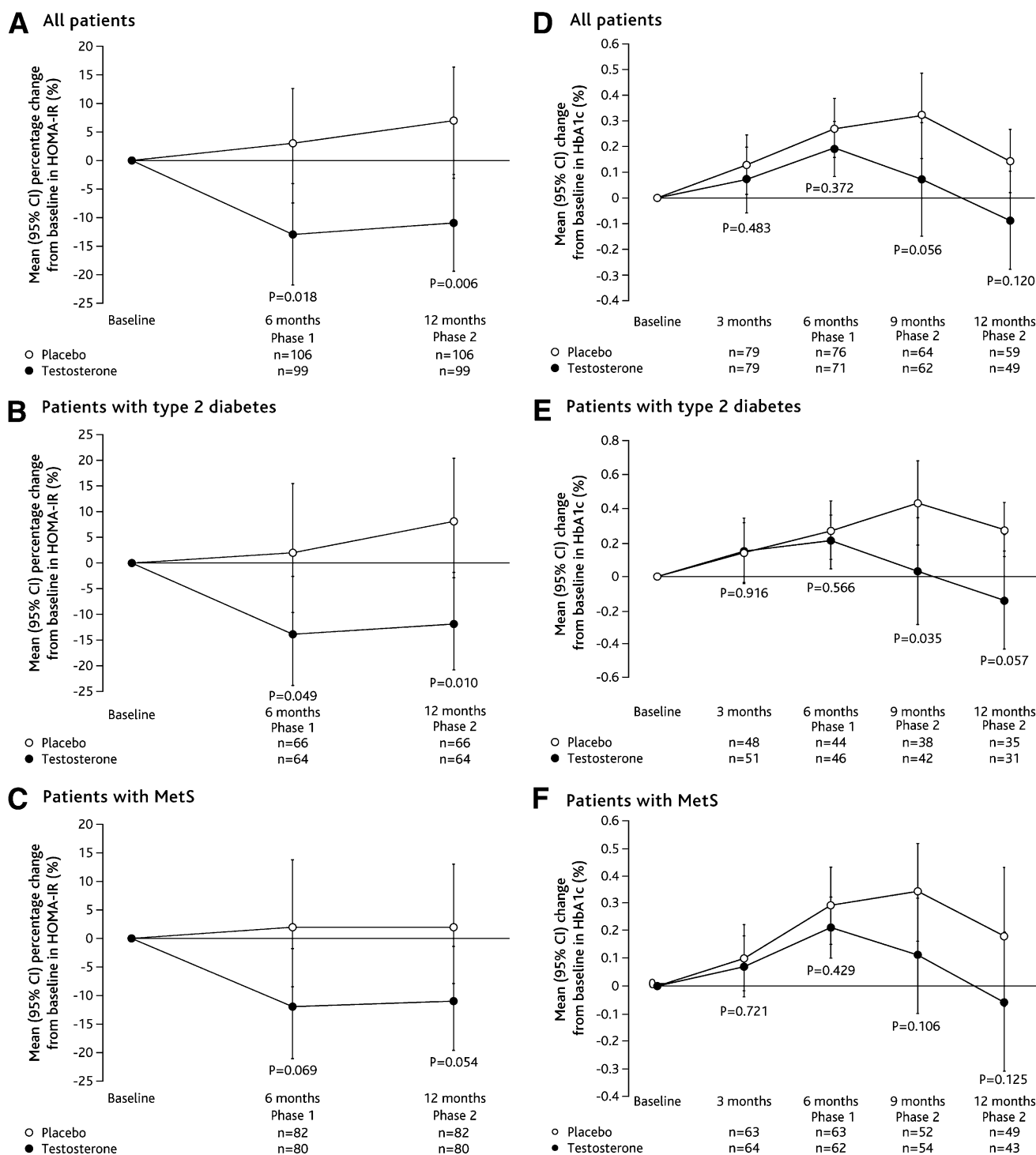


Figure 1—Mean (95% CI) percentage change from baseline in HOMA-IR (ITT population, last observation carried forward) and change from baseline in HbA_{1c} (ITT population, study completers) among all patients (A and D), patients with type 2 diabetes (B and E), and patients with MetS (C and F). P values reported for comparisons between groups.

(TRT -0.92 ± 2.454 , placebo -0.06 ± 2.451 ; $P = 0.17$). However, reductions from baseline in Lpa, TC, LDL cholesterol, and HDL cholesterol were significantly greater in the TRT group than in the placebo group (overall PP cohort). In the exploratory mPP population, TRT was

associated with significant reductions from baseline in percentage body fat, Lpa, TC, and LDL cholesterol in all patients and in the MetS subgroup (Supplementary Table A3). The type 2 diabetes subgroup showed significant reductions in Lpa and percentage body fat

with TRT (Supplementary Table A3). There were no significant between-group differences in HDL cholesterol in the mPP population (Supplementary Table A3). No other end points were significantly different between treatment groups in the PP or mPP populations.

Table 2—Mean values for primary and secondary end points at baseline and months 6 and 12 (ITT population, last observation carried forward)

	All patients		MetS patients		Type 2 diabetic patients	
	Testosterone	Placebo	Testosterone	Placebo	Testosterone	Placebo
Glucose metabolism						
Fasting serum insulin (pmol/L)						
Baseline	151.68 ± 153.26	137.65 ± 147.46	162.17 ± 166.29	153.76 ± 163.02	145.08 ± 158.55	126.19 ± 109.17
n	102	105	82	81	64	65
Phase 1: Month 6	124.59 ± 98.65	128.20 ± 99.76	125.08 ± 86.89	135.50 ± 104.45	121.61 ± 102.01	136.40 ± 117.44
n	101	101	83	80	65	63
Phase 2: Month 12	134.25 ± 108.57	126.89 ± 94.31	135.50 ± 100.92	131.33 ± 98.47	130.91 ± 113.99	131.96 ± 104.30
n	103	101	85	80	65	63
FPG (mmol/L)						
Baseline	7.94 ± 2.99	7.59 ± 2.62	7.60 ± 2.41	7.60 ± 2.71	9.05 ± 3.18	8.49 ± 2.84
n	104	107	84	83	67	67
Phase 1: Month 6	7.84 ± 3.03	7.99 ± 2.86	7.67 ± 2.82	7.88 ± 2.99	8.91 ± 3.31	9.09 ± 2.97
n	102	101	84	80	65	63
Phase 2: Month 12	7.97 ± 3.45	8.14 ± 3.19	7.74 ± 3.25	8.05 ± 3.41	9.18 ± 3.75	9.35 ± 3.46
n	102	101	84	80	65	63
HOMA-B†						
Baseline	104.8 ± 77.5	99.9 ± 75.8	110.5 ± 78.3	104.7 ± 74.7	83.3 ± 67.0	84.1 ± 74.0
n	99	105	80	81	64	66
Phase 1: Month 6	100.7 ± 78.9	98.3 ± 72.7	103.9 ± 79.4	106.0 ± 74.9	79.4 ± 66.0	75.3 ± 58.2
n	105	108	86	84	67	67
Phase 2: Month 12	102.0 ± 79.5	99.4 ± 70.8	105.1 ± 80.1	105.4 ± 70.2	79.1 ± 65.4	79.3 ± 66.5
n	105	109	86	85	67	68
Lipid parameters						
Lpa (μmol/L)						
Baseline	1.44 ± 1.56	1.40 ± 1.17	1.50 ± 1.64	1.42 ± 1.21	1.36 ± 1.36	1.37 ± 1.28
n	98*	102	81**	79	62	64
Phase 1: Month 6	1.22 ± 1.19	1.59 ± 1.80	1.22 ± 1.20	1.49 ± 1.65	1.20 ± 1.16	1.61 ± 2.15
n	96*	98	80**	79	62	60
Phase 2: Month 9	1.39 ± 1.39	1.90 ± 1.94	1.37 ± 1.43	1.77 ± 1.58	1.51 ± 1.52	1.96 ± 2.34
n	51*	56	43	44	35	35
Phase 2: Month 12	1.26 ± 1.22	1.62 ± 1.82	1.26 ± 1.24	1.49 ± 1.38	1.29 ± 1.26	1.65 ± 2.19
n	99	100	83**	80	63	61
TC (mmol/L)						
Baseline	4.65 ± 1.18	4.87 ± 1.11	4.79 ± 1.22	4.98 ± 1.09	4.51 ± 1.17	4.55 ± 1.01
n	106	108	86	84	67	67
Phase 1: Month 6	4.52 ± 1.11	4.89 ± 1.13	4.56 ± 1.11	5.05 ± 1.13	4.44 ± 1.12	4.57 ± 0.97
n	103	102	85**	81	66	63
Phase 2: Month 9	4.55 ± 1.24	4.80 ± 1.12	4.61 ± 1.24	4.95 ± 1.10	4.32 ± 1.14	4.59 ± 1.13
n	62	65	54	52	42	39
Phase 2: Month 12	4.49 ± 1.16	4.77 ± 1.07	4.52 ± 1.17	4.88 ± 1.05	4.31 ± 1.05	4.52 ± 0.95
n	104	102	86	81	66	63
LDL cholesterol (mmol/L)						
Baseline	2.78 ± 0.99	2.89 ± 0.96	2.89 ± 1.02	2.98 ± 0.95	2.61 ± 0.91	2.61 ± 0.82
n	106	108	86	84	67	67
Phase 1: Month 6	2.65 ± 0.91	2.85 ± 0.91	2.70 ± 0.95	2.99 ± 0.93	2.60 ± 0.85	2.55 ± 0.70
n	103	102	85*	81	66	63
Phase 2: Month 9	2.67 ± 1.01	2.77 ± 0.88	2.70 ± 1.01	2.90 ± 0.90	2.53 ± 0.93	2.51 ± 0.77
n	62	65	54	52	42	39
Phase 2: Month 12	2.59 ± 0.91	2.75 ± 0.89	2.62 ± 0.93	2.85 ± 0.91	2.49 ± 0.82	2.50 ± 0.70
n	104	102	86	81	66	63
HDL cholesterol (mmol/L)						
Baseline	1.19 ± 0.30	1.23 ± 0.28	1.21 ± 0.31	1.21 ± 0.26	1.16 ± 0.31	1.23 ± 0.29
n	106	108	86	84	67	67
Phase 1: Month 6	1.12 ± 0.26*	1.21 ± 0.25	1.12 ± 0.24	1.20 ± 0.24	1.09 ± 0.27	1.21 ± 0.26

Table 2—Continued

	All patients		MetS patients		Type 2 diabetic patients	
	Testosterone	Placebo	Testosterone	Placebo	Testosterone	Placebo
<i>n</i>	103	102	85*	81	66*	63
Phase 2: Month 9	1.14 ± 0.31	1.20 ± 0.27	1.14 ± 0.28	1.18 ± 0.27	1.12 ± 0.33	1.20 ± 0.27
<i>n</i>	62	65	54*	52	42	39
Phase 2: Month 12	1.14 ± 0.30	1.21 ± 0.26	1.14 ± 0.28	1.20 ± 0.25	1.09 ± 0.30	1.21 ± 0.27
<i>n</i>	104	102	86	81	66	63
Triglycerides (mmol/L)						
Baseline	1.92 ± 1.36	2.10 ± 1.44	1.94 ± 1.42	2.21 ± 1.52	2.06 ± 1.58	2.05 ± 1.30
<i>n</i>	106	108	86	84	67	67
Phase 1: Month 6	1.93 ± 1.23	2.04 ± 1.28	1.92 ± 1.05	2.10 ± 1.34	1.98 ± 1.31	2.07 ± 1.30
<i>n</i>	103	102	85	81	66	63
Phase 2: Month 9	2.00 ± 1.21	2.23 ± 1.71	2.07 ± 1.26	2.29 ± 1.76	1.91 ± 0.98	2.36 ± 1.99
<i>n</i>	62	65	54	52	42	39
Phase 2: Month 12	1.90 ± 1.15	2.14 ± 1.49	1.91 ± 1.06	2.16 ± 1.54	1.90 ± 1.17	2.19 ± 1.49
<i>n</i>	104	102	86	81	66	63
Abdominal obesity and body composition						
Body fat (%)						
Baseline	33.60 ± 6.54	32.24 ± 6.33	33.99 ± 6.69	32.37 ± 6.17	33.58 ± 5.43	32.54 ± 6.55
<i>n</i>	108	112	88	88	68	69
Phase 1: Month 6	33.01 ± 6.83	32.31 ± 5.77	33.31 ± 6.82	32.43 ± 5.59	32.86 ± 5.92	32.33 ± 5.88
<i>n</i>	94	98	78	78	61	62
Phase 2: Month 12	32.81 ± 7.08	32.63 ± 5.57	33.08 ± 7.18	32.66 ± 5.51	32.69 ± 6.44	32.68 ± 5.71
<i>n</i>	100	98	83	78	63	62
BMI (kg/m ²)						
Baseline	32.87 ± 6.58	31.29 ± 5.44	33.27 ± 6.80	31.63 ± 5.07	32.76 ± 6.12	31.56 ± 5.87
<i>n</i>	108	112	88	88	68	69
Phase 1: Month 6	33.07 ± 6.96	31.45 ± 5.26	33.28 ± 7.28	31.67 ± 5.03	33.16 ± 6.35	31.70 ± 5.55
<i>n</i>	97	99	80	79	62	62
Phase 2: Month 12	32.90 ± 7.01	31.43 ± 5.27	33.17 ± 7.35	31.53 ± 5.07	32.92 ± 6.52	31.73 ± 5.57
<i>n</i>	102	99	84	79	64	62
Waist circumference (cm)						
Baseline	112.7 ± 13.22	110.1 ± 13.77	113.3 ± 13.43	110.8 ± 12.65	112.7 ± 13.35	111.7 ± 15.23
<i>n</i>	108	112	88	88	68	69
Phase 1: Month 6	111.9 ± 13.64	110.2 ± 13.36	112.0 ± 13.62	110.3 ± 12.02	113.1 ± 13.57	111.8 ± 14.46
<i>n</i>	96	99	80	79	61	62
Phase 2: Month 12	111.6 ± 13.89	110.6 ± 13.96	112.0 ± 13.92	110.8 ± 12.52	111.9 ± 13.79	112.4 ± 15.52
<i>n</i>	100	99	83	79	63*	62
Blood pressure						
Systolic blood pressure (mmHg)						
Baseline	138.6 ± 17.30	136.7 ± 17.12	—	—	—	—
<i>n</i>	108	112	—	—	—	—
Phase 2: Month 12	138.7 ± 15.15	134.9 ± 16.49	—	—	—	—
<i>n</i>	50	58	—	—	—	—
Diastolic blood pressure (mmHg)						
Baseline	82.5 ± 10.23	81.6 ± 9.50	—	—	—	—
<i>n</i>	108	112	—	—	—	—
Phase 2: Month 12	82.8 ± 9.72	80.3 ± 9.80	—	—	—	—
<i>n</i>	50	58	—	—	—	—
Sexual dysfunction						
IIEF total score						
Baseline	32.4 ± 18.47	32.8 ± 21.24	—	—	—	—
<i>n</i>	94	98	—	—	—	—
Phase 1: Month 6	41.3 ± 19.97	39.3 ± 22.12	—	—	—	—
<i>n</i>	82*	81	—	—	—	—

Table 2—Continued

	All patients		MetS patients		Type 2 diabetic patients	
	Testosterone	Placebo	Testosterone	Placebo	Testosterone	Placebo
Phase 2: Month 12	40.9 ± 20.34	37.3 ± 22.08	—	—	—	—
n	88*	88				
Erectile function domain score						
Baseline	12.0 ± 9.07	11.9 ± 10.02				
n	101	103				
Phase 1: Month 6	15.6 ± 9.53	15.1 ± 10.49	—	—	—	—
n	84	89				
Phase 2: Month 12	15.2 ± 9.71	14.4 ± 10.42	—	—	—	—
n	90	93				
Orgasmic function domain score						
Baseline	4.6 ± 3.80	4.7 ± 4.20	—	—	—	—
n	105	107				
Phase 1: Month 6	5.9 ± 3.89	5.8 ± 4.18	—	—	—	—
n	88	90				
Phase 2: Month 12	5.5 ± 3.91	5.3 ± 4.23	—	—	—	—
n	95	93				
Sexual desire domain score						
Baseline	5.3 ± 2.13	5.6 ± 2.47	—	—	—	—
n	105	106				
Phase 1: Month 6	6.5 ± 2.27	5.9 ± 2.38	—	—	—	—
n	89**	91				
Phase 2: Month 12	6.3 ± 2.21	5.8 ± 2.55	—	—	—	—
n	94**	95				
Intercourse satisfaction domain score						
Baseline	4.7 ± 4.58	4.7 ± 4.84	—	—	—	—
n	106	108				
Phase 1: Month 6	6.2 ± 4.95	6.3 ± 4.99	—	—	—	—
n	88	91				
Phase 2: Month 12	6.4 ± 5.01	5.7 ± 4.89	—	—	—	—
n	94*	92				
Overall sexual satisfaction domain score						
Baseline	4.7 ± 2.46	4.8 ± 2.68	—	—	—	—
n	96	99				
Phase 1: Month 6	6.0 ± 2.62	5.7 ± 2.81	—	—	—	—
n	85	89				
Phase 2: Month 12	5.9 ± 2.70	5.5 ± 2.73	—	—	—	—
n	91	94				
AMS total score						
Baseline	40.6 ± 11.44	39.9 ± 11.46	—	—	—	—
n	88	96				
Phase 1: Month 6	37.1 ± 11.72	36.1 ± 10.60	—	—	—	—
n	80	86				
Phase 2: Month 12	36.4 ± 11.16	37.8 ± 11.57	—	—	—	—
n	89	92				

Data are means ± SD or n. **P* < 0.05. ***P* < 0.01 between groups. †HOMA-B = (20 × FPI [mIU/L]) / (FPG [mmol/L] − 3.5) (calculated excluding FPG values ≤ 3.5 mmol/L [formula invalid] and atypical insulin values [≥249 pmol/L screening values substituted when baseline values unavailable]).

Efficacy: Phase 2

At the end of Phase 2 (12 months), serum TT values in the ITT population were significantly higher following TRT (19.5 ± 17.0 nmol/L) versus placebo (−0.6 ± 3.3 nmol/L) (*P* < 0.001).

Changes in concomitant medications were allowed in Phase 2. Efficacy results are therefore not presented for this phase

except for HOMA-IR (the primary end point) and sexual function, a measure of hypogonadism. In the total ITT population, HOMA-IR was significantly improved with TRT versus placebo at 12 months (TD 16.4% [95% CI 5–26]; *P* = 0.006).

Exploratory analysis of the type 2 diabetes mPP subgroup showed HbA_{1c}

decreased slightly by month 9 with TRT but increased with placebo, creating a difference of −0.58% that was significant (*P* = 0.005). This trend continued nonsignificantly at 12 months (−0.49%, *P* = 0.066).

There was a significant improvement in IIEF total score and sexual desire and intercourse satisfaction domain scores in

all patients at 12 months in the ITT population (Table 2). The positive effect of TRT on the erectile function domain in the ITT population (Table 2) approached significance in the mPP exploratory analysis (TD 3.225 [95% CI -0.454 to 6.904]; $P = 0.089$).

Safety: Phases 1 and 2

In the safety population, 137 patients (62%) experienced 502 AEs with 224 (45%) considered to be related to study medication (TRT, 116; placebo, 108). The majority of AEs were classified as mild or moderate (TRT, 98%; placebo, 97%) with no significant differences between treatments. The most common AEs in patients experiencing ≥ 1 AE were erythema (TRT, 4%; placebo, 5%), pruritis (TRT, 4%; placebo, 4%), and nasopharyngitis (TRT, 3%; placebo 4%) (Supplementary Table A5).

A total of 17 serious AEs were experienced by 13 patients. Of these, 3 in the placebo group were considered related to treatment. Thirteen resolved without sequelae. One patient (placebo) died as the result of myocardial infarction. A total of 30 patients withdrew from the study because of AEs (TRT, 18; placebo, 12). The most common AEs leading to discontinuation were skin and subcutaneous tissue disorders (21 AEs; testosterone, 11; placebo, 10). Cardiovascular events occurred more commonly with placebo (10.7 vs. 4.6%; $P = 0.095$).

No clinically relevant mean changes in laboratory parameters were noted. After 12 months, hematocrit had increased by 0.03 ± 0.04 L/L for TRT versus a decrease of -0.01 ± 0.02 L/L for placebo, and hemoglobin levels had increased by 1.42 ± 1.55 g/dL from baseline (TRT) versus 0.06 ± 0.7 g/dL (placebo). There was no significant increase in age-adjusted PSA values. The PSA level exceeded the upper limit of age-adjusted normal values in three subjects at baseline (testosterone, 2; placebo, 1). This violation of the entry criteria would not have affected the primary efficacy end point, and because the patients had been randomized and were satisfactorily proceeding in the study they were not withdrawn. None of these patients underwent prostate biopsy during the study. PSA levels exceeded normal limits in four subjects at 12 months (TRT, 3; placebo, 1).

CONCLUSIONS—This study demonstrates that transdermal TRT in hypogonadal

men with type 2 diabetes and/or MetS improves IR, the central biochemical defect associated with these conditions. This result is of added interest, as many study participants were already taking OADs (mainly metformin), which affect insulin sensitivity. Although an improvement in IR could be expected to result in better glycemic control, no reduction in HbA_{1c} was demonstrated after 6 months of treatment. Previous studies have reported that TRT does improve glycemic control (6,9); however, the current study was not powered to detect changes in glycemic control and included patients with controlled diabetes (HbA_{1c} <6.5%). A significant difference in HbA_{1c} between the TRT and placebo groups was observed at 9 months, but this potential treatment effect was confounded by the permitted changes in diabetes-related medications for ethical reasons. Therefore, no clear conclusion can be made. A larger definitive study with HbA_{1c} as a primary endpoint in hypogonadal men with uncontrolled diabetes is required to investigate this further.

The majority of cross-sectional studies have shown that hypogonadism is associated with dyslipidemia, albeit with conflicting results (4). There are only limited data supporting the effect of testosterone gel formulations on lipids. Two studies reported a small rise in HDL cholesterol but no effect on TC or LDL cholesterol in hypogonadal men (9,21). In the current study, Lpa was significantly reduced compared with placebo in the overall population after 6 months of therapy. Moreover, the subgroup of patients with MetS at baseline showed significant reductions from baseline with TRT versus placebo, in Lpa, TC, and LDL cholesterol after 6 months, even though the majority of these patients were taking statins and had mean baseline TC concentrations below 5 mmol/L. A reduction in HDL cholesterol has been reported in studies using intramuscular testosterone ester therapy (4).

Lpa is a strong independent risk factor for premature coronary heart disease (22). Surgical orchidectomy for prostate cancer increases Lpa concentrations, whereas administration of testosterone in healthy men reduces Lpa levels (23). In the current study, reductions from baseline in serum Lpa concentration were significantly greater in TRT-treated patients at month 6 in both the overall population and the MetS subgroup. The long-term

cardiovascular benefits of reducing Lpa are unknown.

The safety and tolerability profile of TRT was comparable to placebo with no significant differences in cardiovascular events, including blood pressure. Caution has been urged following a recent study that reported an increase in cardiovascular events in testosterone-treated, frail, elderly men (24). However, cardiovascular events were not a planned outcome in this study, and verification of events varied among subjects. Moreover, >70% of the men in that trial received higher doses of testosterone (100 or 150 mg/day) than any of the men in the current study (40–80 mg/day). The current study showed a trend for cardiovascular events to occur more frequently in the placebo group than in the TRT group. A similar frailty study that used standard doses (testosterone gel 50 mg/day) in clinical practice found no increase in cardiovascular events (25).

The frequency of application site reactions in this study was higher than has been reported in other trials involving transdermal TRT. However, in TIMES2 mandatory application site inspections were performed at each study visit, and the investigators were trained to identify and document skin-related reactions. This is in contrast to other studies that relied on patients reporting a dermatological AE.

The beneficial effects of TRT on overall sexual function and, importantly, libido (a key symptom of testosterone deficiency) support the assessment and treatment of hypogonadism in men with type 2 diabetes and/or MetS who do not present specifically with erectile dysfunction.

Results from the mPP population should be regarded as exploratory only, since this population was created as part of a post hoc analysis. However, these results have been included because they allow assessment of the impact of testosterone on individual parameters without the potential confounding influence of changes in concomitant medication. Although this information is typically gained from the PP population, the small size of the PP population here limited the power of any analyses and precluded analysis of disease subgroup data (MetS and type 2 diabetes). The mPP data provide supportive evidence that TRT improves glycemic control, lipid profile, central obesity and body composition in hypogonadal men with type 2 diabetes

and/or MetS. In this population, the loss of statistical significance for some end points between 6 and 12 months is likely to be due to the smaller size of the population at month 12 and the allowed use of concomitant medications after 6 months (Supplementary Table A3). The delayed effect of TRT on HbA_{1c} could be explained by the cumulative effect on reduction in body fat over several months.

The high dropout rate is a limitation of this study. However, there was no difference between treatment groups in the number or characteristics of the patients who failed to complete the study.

This study has shown that TRT in hypogonadal men with type 2 diabetes and/or MetS improves several cardiovascular risk factors, most notably IR, Lp(a), TC, and LDL cholesterol. These findings and the therapeutic benefit on sexual function strongly support a role for TRT in these clinical conditions.

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T.H.J. and K.S.C. conceived and designed the study protocol and produced the initial draft of the manuscript. T.H.J., K.S.C., S.A.,

H.M.B., J.B., E.M., I.M., A.M.M., M.V., A.Y., and J.D.H. contributed to data analysis and interpretation and to the writing and editing of the manuscript. All authors had access to all data and had final responsibility for the manuscript content; T.H.J. had final responsibility for the decision to submit for publication.

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