



Office de la Propriété
Intellectuelle
du Canada

Un organisme
d'Industrie Canada

Canadian
Intellectual Property
Office

An agency of
Industry Canada

CA 2453337 C 2012/08/28

(11)(21) **2 453 337**

(12) **BREVET CANADIEN
CANADIAN PATENT**

(13) **C**

(86) Date de dépôt PCT/PCT Filing Date: 2002/07/09
(87) Date publication PCT/PCT Publication Date: 2003/01/23
(45) Date de délivrance/Issue Date: 2012/08/28
(85) Entrée phase nationale/National Entry: 2004/01/08
(86) N° demande PCT/PCT Application No.: US 2002/021524
(87) N° publication PCT/PCT Publication No.: 2003/005954
(30) Priorité/Priority: 2001/07/09 (US60/304,313)

(51) Cl.Int./Int.Cl. *A61K 31/138* (2006.01),
A61K 9/48 (2006.01), *A61P 5/28* (2006.01)
(72) Inventeur/Inventor:
PODOLSKI, JOSEPH S., US
(73) Propriétaire/Owner:
REPROS THERAPEUTICS INC., US
(74) Agent: RIDOUT & MAYBEE LLP

(54) Titre : METHODES ET MATERIAUX DESTINES AU TRAITEMENT DE LA DEFICIENCE EN TESTOSTERONE
CHEZ L'HOMME

(54) Title: METHODS AND MATERIALS FOR THE TREATMENT OF TESTOSTERONE DEFICIENCY IN MEN

(57) **Abrégé/Abstract:**

The present invention relates to the use of compositions comprising trans-clomiphene for treating men with hypogonadism. The invention is also directed to methods for treating males with hypogonadism.



(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
23 January 2003 (23.01.2003)

PCT

(10) International Publication Number
WO 03/005954 A3

(51) International Patent Classification⁷: **A61K 31/225**,
31/135

(21) International Application Number: PCT/US02/21524

(22) International Filing Date: 9 July 2002 (09.07.2002)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/304,313 9 July 2001 (09.07.2001) US

(71) Applicant (for all designated States except US): **ZONA-
GEN, INC.** [US/US]; 2408 Timberloch Place, B-4, The
Woodlands, TX 77380 (US).

(72) Inventor; and

(75) Inventor/Applicant (for US only): **PODOLSKI, Joseph,**
S. [US/US]; 3 Pebble Hollow Court, The Woodlands, TX
77381 (US).

(74) Agent: **CLOUGH, David, W.**; Katten Muchin Zavis
Rosenman, 525 West Monroe Street, Suite 1600, Chicago,
IL 60661 (US).

(81) Designated States (national): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,

CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,
MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG,
SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
VN, YU, ZA, ZM, ZW.

(84) Designated States (regional): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),
Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),
European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE,
ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK,
TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ,
GW, ML, MR, NE, SN, TD, TG).

Published:
— with international search report
— with amended claims and statement

(88) Date of publication of the international search report:
23 October 2003

Date of publication of the amended claims and statement:
4 December 2003

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

(54) Title: METHODS AND MATERIALS FOR THE TREATMENT OF TESTOSTERONE DEFICIENCY IN MEN

(57) Abstract: The present invention relates to the use of compositions comprising *trans*-clomiphene for treating men with hypogonadism. The invention is also directed to methods for treating males with hypogonadism.



WO 03/005954 A3

METHODS AND MATERIALS FOR THE TREATMENT OF TESTOSTERONE
DEFICIENCY IN MEN

[0001]

FIELD OF THE INVENTION

[0002] The present invention relates to the compositions and methods for increasing testosterone levels. More specifically, the present invention relates to a composition comprising clomiphene enriched for *trans*-clomiphene. The present invention also relates to the use of a composition comprising clomiphene enriched for *trans*-clomiphene reagents for increasing testosterone levels.

BACKGROUND

[0003] Testosterone is the primary male androgen, playing a vital role in overall male health. Testosterone is essential to the development and maintenance of specific reproductive tissues (testes, prostate, epididymis, seminal vesicle, and penis) and male secondary sex characteristics. It plays a key role in libido and erectile function and is necessary for the initiation and maintenance of spermatogenesis. Testosterone also has important functions not related to reproductive tissues. For example, it positively affects body composition by increasing nitrogen retention, which supports lean body mass, muscle size and strength. It also acts on bone to stimulate bone formation.

[0004] Testosterone secretion is the end product of a series of hormonal processes. Gonadotropin-releasing hormone (GnRH), which is secreted in the hypothalamus, controls the pulsatile secretion of luteinizing hormone (LH) and follicle stimulating hormone (FSH), which are secreted by the anterior pituitary. LH, in turn, regulates the production and

secretion of testosterone in the Leydig cells of the testes, while FSH assists in inducing spermatogenesis.

[0005] Testosterone is most often measured as "total testosterone." This measurement includes testosterone that is bound to sex hormone-binding globulin (SHBG) (~44%) and is therefore not bioavailable and testosterone which either is free (~2%) or loosely bound to other proteins (non-SHBG-bound) (~54%).

[0006] Results from a WHO study indicate that testosterone is normally secreted in a circadian rhythm, with higher levels in the morning and nadir levels occurring around 8 to 10 p. m. *See FIG. 1.* This variation in testosterone secretion throughout the day becomes much less pronounced in older men (mean age equals 71 years). The importance of this rhythm is not known at this time.

[0007] Samples were obtained from both young and elderly patients every 10 minutes for 24 hours via an indwelling cannula. According to Tenover J. *et al.*, (Journal Clinical Endocrinol. Metab. Vol. 65, No. 6, pp. 1118-1126 (1987)) the mean 24 hr total serum testosterone levels in healthy young men (age range 22 yrs.-35 yrs. mean 27.3 yrs) was 4.9 ± 0.3 (\pm SEM) mg/ml (17.0 nmol/L) while older men (age range 65yrs-84 yrs. mean 70.7 yrs.) had a significantly lower mean 24 hrs. total serum testosterone level of 4.1 ± 0.4 mg/ml. ($P < 0.5$; 14.2 nmol/L).

[0008] Total serum testosterone levels obtained from single random samples were also significantly lower in older men (4.0 ± 0.2 mg/ml [13.9 n nmol/L]) as compared to 4.8 ± 0.2 mg/ml [16.6 nmol/L] in healthy young men.

[0009] Testosterone deficiency can result from underlying disease or genetic disorders and is also frequently a complication of aging. For example, primary hypogonadism results from primary testicular failure. In this situation, testosterone levels are low and levels of pituitary gonadotropins (LH and FSH) are elevated. Secondary hypogonadism is due to inadequate secretion of the pituitary gonadotropins. In addition to a low testosterone level, LH and FSH

levels are low or low-normal. Some of the sequelae of adult testosterone deficiency include a wide variety of symptoms including: loss of libido, erectile dysfunction, oligospermia or azoospermia, absence or regression of secondary sexual characteristics, progressive decrease in muscle mass, fatigue, depressed mood and increased risk of osteoporosis.

[0010] Several forms of testosterone therapy exists in the United States today. Recently, transdermal preparations have gained favor in the market. However, a scrotal testosterone patch results in supraphysiologic levels of 5 α -dihydrotestosterone (DHT) due to the high concentration of 5 α -reductase in scrotal skin. It is not known whether these elevated DHT levels have any long-term health consequences. Nonscrotal systems are considered more convenient and most patients achieve average serum concentrations within the normal range and have normal levels of DHT. Oral testosterone therapy is not recommended because doses required for replacement therapy are associated with significant risk of hepatotoxicity.

SUMMARY

[0011] The present invention is directed to compositions useful for increasing testosterone levels in male mammals and for ameliorating or preventing the sequelae of low testosterone levels. In one of its aspects the invention is directed to compositions having active ingredients comprising 0% to 29% weight/weight of (*cis*, -Z-, *trans*-clomiphene) (hereinafter "*cis*-clomiphene") and 100% to 71% w/w (*trans*-, E-, *cis*-clomiphene) (hereinafter "*trans*-clomiphene") or pharmaceutically acceptable salts thereof. Among the preferred compositions of the present invention which contain both *cis*-clomiphene and *trans*-clomiphene invention are compositions wherein the ratio of *trans*-clomiphene and *cis*-clomiphene is greater than 1. A more preferred composition according to the present invention comprises about 100% w/w of active ingredients of *trans*-clomiphene or a pharmaceutically acceptable salt thereof. All compositions of the present invention may further comprise suitable pharmaceutical excipients diluents, carriers, and the like. Analogs

of the *cis*-clomiphene and *trans*-clomiphene are also contemplated for use in all aspects of the present invention.

[0012] The present invention is also directed to methods for increasing serum testosterone levels in hypogonadal male mammals (and for ameliorating or preventing the sequelae of low testosterone levels), the method comprising administering to a subject male an effective amount of a composition according to the present invention, the compositions having active ingredients comprising 0% to 29% weight/weight of *cis*-clomiphene and 100% to 71% w/w *trans*-clomiphene including any of their pharmaceutically acceptable salts thereof. Among the preferred methods are those in which the administered compositions contain both isomers wherein the ratio of *trans*-clomiphene to *cis*-clomiphene is greater than 1. A more preferred method comprises administering to the male a composition comprising about 100% w/w of *trans*-clomiphene.

BRIEF DESCRIPTION OF THE DRAWING

[0013] FIG. 1 is a graphic representative of the normal secretory total serum testosterone profiles in healthy men (young and old).

[0014] FIG. 2 shows the chemical structure of clomiphene citrate.

[0015] FIG. 3 is a graphic demonstration of the time course of serum testosterone levels with Clomid, Enclomid and Zuclomid.

DETAILED DESCRIPTION

[0016] The present invention provides methods and compositions useful for increasing testosterone levels in male mammals and for ameliorating or preventing the sequelae of low testosterone levels including but not limited to those described above.

[0017] Clomiphene (FIG. 2) is an antiestrogen related to tamoxifen that blocks the normal estrogen feedback on the hypothalamus and subsequent negative feedback on the pituitary. This leads to increases in luteinizing hormone (LH) and follicle stimulating hormone (FSH).

In men, these increased levels of gonadotropins stimulate the Leydig cells of the testes and result in the production of higher testosterone levels. Clomiphene citrate has the following structure:

[0018] Ernst *et al.*, (J. Pharmaceut. Sci. 65 : 148 (1976)), have shown that clomiphene is a mixture of two geometric isomers which they refer to as *cis*-, *Z*-, clomiphene (*cis*-clomiphene or zuclophene) and *trans*-, *E*-, clomiphene, (*trans*-clomiphene or enclophene).

According to Ernst, *et al.*, (J. Pharmaceut. Sci. 65: 148 (1976)), *trans*-clomiphene HCl has a melting point of 149°C-150.5°C, while *cis*-clomiphene HCl has a melting point of 156.5°C-158°C.

[0019] Ernst, *et al.*, (J. Pharmaceut. Sci. 65: 148 (1976)) have also noted that (the *trans*-isomer) is antiestrogenic (AE) while the *cis*-isomer is the more potent and more estrogenic form and has also been reported to have anti-estrogenic activity. The authors attribute the effect of the drug on ovulatory activity to both forms stating that the mixture is more effective than *trans*-clomiphene alone. The *trans*- isomer aids ovulation at the level of the hypothalamus. The estrogenic isomer *cis*- clomiphene contributes to enhanced ovulation elsewhere in the physiologic pathway leading to ovulation. The isomers are also reported to have different *in vivo* half-life. Furthermore the *cis* form has been reported to leave residual blood levels for in excess of one month following a single dose.

[0020] Vandekerckhove, *et al.* (Cochrane Database Syst Rev 2000; (2): CD000151 (2000)) noted that ten studies involving 738 men have suggested that anti-estrogens appear to have a beneficial effect on endocrinal outcomes, *i. e.* testosterone, but there is not enough evidence to evaluate fertility effects. Nevertheless should clomiphene administration enhance testosterone levels then one could easily conclude that the drug should positively impact the side effects of testosterone deprivation as long as the testes still retain the ability to respond to gonadotropin stimulation.

[0021] Clomiphene is currently approved as a mixture of both cis- and trans-isomers, the cis-isomer being present as about 30% to 50% (Merck Manual) for fertility enhancement in the anovulatory patient. Clomiphene improves ovulation by initiating a series of endocrine events culminating in a preovulatory gonadotropin surge and subsequent follicular rupture. The drug is recommended to be administered for 5 days at a dose of up to 100 mg daily. Clomiphene has also been associated with numerous side effects including: blurred vision, abdominal discomfort, gynecomastia, testicular tumors, vasomotor flushes, nausea, and headaches. Furthermore, other studies suggest that clomiphene possesses both genotoxic and tumor enhancement effects. The net outcome of these observations is that clomiphene in its current format, having between 30% and 50% of the cis isomer, would be unacceptable for chronic therapy in men for the treatment of testosterone deficiency.

[0022] Clomiphene has also been used for therapeutic intervention in men with low testosterone levels. Tenover *et al.*, J. Clin. Endocrinol. Metab. 64:1103, (1987) and Tenover *et al.*, J. Clin. Endocrinol. Metab. 64:1118 (1987) found increased in FSH, LH in both young and old men after treatment with clomiphene. They also found increases in free and total testosterone in men with young men showing significant increases

[0023] Studies were also conducted to determine whether or not clomiphene could be used to improve fertility in men by improving semen quality. Homonnai *et al.* Fertil. and Steril 50:801 (1988) saw increases in sperm concentration and count but others have not. (See e.g., Sokel, *et al.*, Fertil. and Steril. 49:865 (1988); Check, *et al.*, Int. J. Fertil. 34:120 (1989); Purvis, *et al.*, Int. J. Androl 21:109 (1989); and Breznik, Arch. Androl. 21:109 (1993).) One group saw a deterioration in the percentage of normal sperm with long-term treatment. Shamis, *et al.*, Arch. Androl 21:109 (1991). A WHO study showed no changes in semen quality or fertility after 6 months of treatment. (Anonymous Androl. 15:299 (1992).) A meta-analysis seems to confirm that testosterone levels go up in men with poor quality sperm

but not fertility. (Vanderkerckhove, *et al.*, 2000 *supra*). Studies have also suggested that long term treatment with clomiphene does not seem to have a drastic deleterious effect on health, although it did show that treatment resulted in poorer sperm quality after 4 months. Studies have kept men on clomiphene for as long as 18 months and at levels of 25 mg per day or 100 mg every other day.

[0024] In 1991, *Guay et al* (Urology 38 : 377 (1991)) suggested that clomiphene could treat sexual dysfunction in men. Their hypothesis seems to be that sexual function follows testosterone levels. This was supported by early studies showing positive influence of androgens and sexual function, Davidson, *et al.*, J. Clin. Endocrinol. Metab. 48 : 955 (1979), and studies that rated sleep-related erections as a strong response to T, Cuxuingham, *et al.*, J. Clin. Endocrinol. Metab. 70: 792 (1990). However, in 1995, *Guay et al.* (Gray, *et al.*, J. Clin. Endocrinol. Metab. 80 : 3546 (1995)) published a study in which they saw increase in LH, FSH, and testosterone after 2 months of clomiphene but no effects on erectile dysfunction.

There might be some advantage for young men and specific groups of older men, but it seems that just raising the testosterone level is not enough. Effects of testosterone on sleep-related erections may have been taken too seriously (Herskowitz, *et al.*, J. Psychosomat. Res. 42 : 541 (1997)).

[0025] According to the present invention, a composition comprising of one isomer preferably *trans*-clomiphene or a redefined blend of the isomers of clomiphene as described below differing from the normally produced mixture are used to enhance testosterone levels while reducing the side effects of the drug. Thus, the present invention provides an oral therapy for increasing testosterone levels, which lacks or has diminished side effects connected with the existing clomiphene formulations.

[0026] In one embodiment of the present invention, a patient who has a need or desire to increase their serum testosterone levels are administered one or more dosages of an effective

amount of composition comprising *trans*-clomiphene at a dosage between one mg to about 200 mg (although the determination of optimal dosages is with the level of ordinary skill in the art). *Cis*-clomiphene may also be present in the composition so long as the ratio of *trans*-clomiphene to *cis*-clomiphene is greater than 1. Analogs of the *trans*- and *cis*-isomers of clomiphene such as those described in Ernst, *et al.*, J. Pharmaceut. Sc. 65: 148 (1976), *supra* are also useful in the practice of the present invention.

[0027] Dosages are preferably (but not necessarily) administered as part of a dosage regimen designed to give rise to serum testosterone levels that mimic or correspond to the normal secretory total serum testosterone profile described in FIG. 1. For example, according to FIG. 1 a dosage of the preferred composition may be administered in a pharmaceutical formulation that would give rise to peak serum testosterone levels at around 8 a. m. Such pharmaceutical formulations may be in the form of sustained release formulations prepared as described for example in U. S. Patent No. 6,221, 399, Japanese patent 4-312522, Meshali *et al.*, Int. J. Phar. 89: 177-181 (1993), Kharenko *et al.*, Intern. Symp. Control Rel. Bioact. Mater.

22: 232-233 (1995), WO 95/35093, Dangprasit *et al.*, Drug. Devel. and Incl. Pharm. 21 (20): 2323-2337 (1995); U. S. Patent Nos. 6,143, 353,6, 190,591, 6,096, 338,6, 129,933, 6,126, 969,6, 248,363 and other sustained release formulations well known in the art.

[0028] Suitable pharmaceutical compositions or unit dosage form may be in the form of solids, such as tablets or filled capsules or liquids such as solutions suspensions, emulsions, elixirs or capsules filled with the same, all for oral use. The compositions may also be in the form of sterile injectable solutions or emulsions for parenteral (including subcutaneous) use.

Such pharmaceutical compositions and unit dosage forms thereof may comprise ingredients in conventional proportions.

[0029] Compositions according to the present invention may also be administered by the intravenous, subcutaneous, buccal, transmucosal, intrathecal, intradermal, intracisternal or

other routes of administration. After administration of the composition serum testosterone levels may be measured as described above and dosages may be altered to achieve a sufficient increase in the serum testosterone levels to achieve the desired physiological results associated with normal testosterone described above.

[0030]

[0031] The following Example is meant to be illustrative of the invention and is not intended to limit the scope of the invention as set out in the appended claims.

EXAMPLE 1

Effects of Clomids on Serum Testosterone in Male Baboons

[0032] Adult, male, Baboons were given 1.5 mg/kg of Clomid, Enclomid (*trans*-Clomid) or Zuclomid (*cis*-Clomid) for 12 consecutive days. The samples analyzed were sera taken on the day of first treatment before being given test article (day 0), after 12 days of treatment (day 12) and 7 days after the last treatment (end or wash-out).

1. Effects on Body Weight and Serum LH, FSH, PRL and Testosterone

[0033] There were significant increases in total serum testosterone in the group receiving Enclomid. See Table 1. There were no differences among groups in the baseline period or at day 0. There were also no differences among the three groups 7 days after treatment (the washout period). However, Enclomid produced higher levels of testosterone compared to Clomid and Zuclomid on day 6 ($p = 0.03$ and $p = 0.00002$ respectively) and compared to Zuclomid on day 12 ($p = 0.047$). Zuclomid clearly did not raise total serum testosterone to any extent. Compared to the animals receiving Enclomid, the animals receiving Clomid exhibited more variable total testosterone levels on day 6 and later as judged by their coefficients of variations. When we looked at the time course of the effects (FIG. 3), we determined that only Enclomid significantly and statistically raised total serum testosterone

on days 6 and 12 compared with either baseline or day 0 values. Moreover, cessation of Enclomid treatment, resulted in a significant drop in the level of total serum testosterone between day 12 and day 18 (washout). This indicates that Enclomid is readily cleared from the circulation consistent with the metabolic clearance seen for Enclomid in humans. Enclomid was clearly better and more consistent than Clomid itself and Zucloamid was ineffective.

Table 1 - Serum Testosterone Levels (ng/dl)

Group	ID	baseline 12/3/01	0 day 12/7/01	6 days 12/13/01	12 days 12/20/01	wash-out 12/26/01
CLO	7500	79.01	76.15	940.97	891.5	150.9
	9012	97.55	305.24	585.92	555.6	316.3
	9097	158.06	102.94	151.12	318.9	143.6
	mean	111.5	161.4	559.3	588.7	203.6
	SD	41.3	125.2	395.6	287.7	97.7
ENCLO	7223	64.57	74.96	1223.8	633.6	307.2
	8021	166.86	133.59	1128.2	1466	399.2
	8369	170.45	106.47	1081.1	1166	271
	mean	134.0	105.0	1144.4	1088.5	325.8
	SD	60.1	29.3	72.7	421.6	66.1
ZUCLO	7438	124.84	210.4	137.51	314.5	359.7
	8292	104.66	67.37	169.98	406.1	860.5
	10098	282.29	904.82	227.95	353.0	274.1
	mean	170.6	394.2	178.5	357.9	498.1
	SD	97.3	448.0	45.8	46.0	316.8
ANOVA		p = 0.61	p = 0.43	p = 0.007	p = 0.57	p = 0.256
K-W		p = 0.56	p = 0.84	p = 0.051	p = 0.079	p = 0.252

[0034] There were no changes in serum LH or FSH. The ratio of total serum testosterone to LH followed the same pattern as total serum testosterone, suggesting a lack of dependence (data not shown). There was also no change in body weight during the 12 day study. There was a decrease in serum prolactin (PRL) during the study in the group receiving Enclomid, suggesting an effect of antiestrogen that has been described in part (Ben-Jonathan and

Hnasko, 2001) and expected on the basis of the fact that as men age, testosterone declines and Prolactin increase (Feldman *et al.*, 2002).

2. Effects on Clinical Chemistry Parameters

[0035] The mean values for each parameter did not differ among the three groups for any test parameter at the beginning of the study as determined by ANOVA or by the Kruskal-Wallis test. All groups exhibited normal values at each parameter except for (1) serum sodium; a related calculated parameter, anionic gap, which were low for all nine baboons throughout the trial; (2) serum glucose; and (3) BUN which were high on day 0 for the group which would be treated with Enclomid. On day 12 of treatment and 7 days after treatment (washout), there were no differences among groups for any parameter except anionic gap that showed that the Clomid and Zuclomid groups had lower values than the Enclomid group. The values of serum sodium and anionic gap appear to be anomalies associated with this group of baboons.

[0036] There were substantive effects on the red blood cell population with Enclomid and Zuclomid and on hematocrit with Zuclomid. All the compounds lower the mean cell hemoglobin concentration (MCHC) either at day 0 or at the endpoint. With no change in mean cell hemoglobin (MCH) and an increase in the mean cell volume (MCV), the lowering of MCHC is predictable. Although testosterone might be expected to raise hematocrit, only Zuclomid treatment, which did not increase total serum testosterone, demonstrated a statistical difference. Clearly, men in a clinical trial that uses Zuclomid should be monitored for the characteristics of their red blood cell population. Enclomid would be predicted to have less of an effect.

[0037] There appears to be a clear effect of 12-day Enclomid treatment on platelets although the values found stayed within the normal range. One thing to consider here is the sexual dimorphism in platelet counts between male and female baboons (279 for males vs. 348 for

females). This is likely to be due to hormones. Since the Enclomid group demonstrated increased testosterone, the lowering of the platelet count could be secondary to the change in testosterone in this group. Moreover, treatment with Enclomid pushed the platelet count to its normal male level from a day 0 level that was the high end of the normal range for this group. Enclomid would not necessarily predict a deleterious effect on platelets.

[0038] All the Clomids tested had effects on the white blood cell (WBC) population, the most striking was that of Enclomid on raising the counts of lymphocytes and eosinophiles. The effects are not as straightforward as they would seem to be. There appears to be a strong effect of Enclonud on lowering the per cent of granulocytes in the blood. The effects are very strong after the 7-day washout period when the values are decreased below the normal range. (This time course could reflect the relatively long time required to affect change the WBC population.) There is little sexual dimorphism in baboons with respect to the white blood cell populations, so the effects are more likely to be due to the compound itself than changes in testosterone. However, when we look at the calculated count of granulocytes using the WBC count, we find no differences in granulocyte count due to any compound. Concomitantly, it is the lymphocyte story that is the most interesting. Both the count and per cent lymphocytes in the population increase with Enclomid treatment. Whereas the mean values of per cent lymphocytes remain in the normal range, given the trend for an increase in WBC count, the net effect is an increase in lymphocyte count with Enclomid. This eosinophil result is analogous. There is a clear implication for treating men who have low lymphocytes, such as men who are HIV-positive. Since Enclomid is unlikely to lower lymphocytes based on this result, a case could be made for its use in the population of men with AIDS. These individuals are often treated with agents that are intended to raise testosterone due to the wasting effects of disease. Low liver and kidney toxicity and favorable effects on cholesterol

and lipids are also highly favored attributes for any medication intended for use HIV-positive men who are already compromised by their disease.

[0039] The increase in serum glucose with Clomid or Zuclomid was within the normal range. In the case of Enclomid where the mean serum glucose values were high on day 0, there were no increases with treatment. There was no evidence that Enclomid would have a deleterious effect on blood glucose.

[0040] No clearly adverse effects on liver function are apparent as judged by the enzymes AST and ALT. The trend in these values was a decrease with treatment. An increase in the level of enzymes in the serum would indicate liver damage. ALT/SGPT was out of range low at the end of the study for the Clomid group although the differences over the treatment period were not statistically significant. The changes with Enclomid and Zuclomid were within the normal range. AST is depressed in pregnancy; thus the action of an estrogen agonist such as Zuclomid in lowering the marginal AST level could be rationalized. Alkaline phosphatase (ALP) is also found in the liver and is elevated various disease states. The lowering of ALP argues further against hepatic damage. There were no changes in serum albumin, also a liver product. A strong suppression of serum albumin over an extended time period could contribute to free serum steroid hormone levels in humans although a more important role is played by sex hormone binding globulin. As a bottom line, none of the compounds could be linked to liver damage on the basis of the parameters assayed.

[0041] Osteoblastic activity and diseases of the bone are accompanied by high serum ALP values. ALP was not elevated following Zuclomid treatment and was decreased in value following Enclomid treatment. The trends would predict a more benign result for the use of Enclomid compared to Zuclomid.

[0042] Although BUN and BUN/creatinine were altered during the study in the Clomid and Enclomid groups, the lack of a definitive change in creatinine argues against renal

dysfunction. A loss of glomerular filtration capacity would result in an increase in BUN.

Decreased BUN occurs in humans due to poor nutrition (not likely in a controlled setting), or high fluid intake (presumably accompanied by edema). Also, despite an increase in total serum testosterone between day 0 and Day 12 with Enclomid, there were no differences between serum creatinine values, arguing against an increase in muscle mass over this short time interval.

[0043] Serum sodium levels were lower than reference values for all animals throughout the study. Serum carbon dioxide was higher than reference values on day 12 for the Clomid and Zuclomid groups. Serum anion gap was lower for all animals throughout the study, paralleling the sodium results. Enclomid raised this parameter towards normal values. The electrolyte imbalances detected in the test animals throughout all treatment periods remains elusive but might be part of the same fluid derangement phenomenon suggested by the BUN results.

[0044] Treatment with Enclomid tended to decrease serum cholesterol and Zuclomid tended to increase the same parameter although neither change reached statistical significance. Those changes were within the normal range although the trend for the two isomers to demonstrate opposite effects over a short period of time merits the further monitoring and might not be unexpected given that the isomers have, alternatively, estrogen agonist or antagonist activity. Enclomid might be expected to be more benign than Zuclomid with respect to serum cholesterol if used chronically.

[0045] The foregoing results indicate that Enclomid is more effective than Clomid or Zuclomid at enhancing total serum testosterone. Zuclomid is clearly not effective and that deficiency limits any use of Clomid for hypogonadism, particularly since the Zuclomid component of Clomid would predominate in the circulation over time given its longer half-life.

[0046] Enclomid appeared to be relatively benign in all aspects when compared to Zuclomid and, often, even Clomid. This is particularly true when consideration is given to the trend of Enclomid to lower cholesterol, and liver enzymes as opposed to Zuclomid's trend to raise the same parameters. The surprising trend for Enclomid to raise the lymphocyte count may be useful for men with AIDS if it can be shown the CD4+ subpopulation of lymphocytes is not lowered or is enhanced.

EXAMPLE 2

Method for Increasing Testosterone Level in Men Using *Trans*-clomiphene and Mixtures of *Trans*-clomiphene and *Cis*-clomiphene at Ratios Greater Than 1

[0047] Prior to administration of *trans*-clomiphene, blood samples are taken from subject males and testosterone levels are measured using methodologies described for example in Matsumoto, *et al.* Clin. Endocrinol. Metab. 56; 720 (1983).

Sex hormone binding globulin (SHBG), both free and bound to testosterone, may also be measured as described for example in Tenover *et al.* J. Clin. Endocrinol. Metab. 65:1118 (1987) which describe measurement of SHBG by both a [³H] dihydrotestosterone saturation analysis and by radioimmunoassay. Non-SHBG-bound testosterone levels (bioavailable testosterone) are also measured for example according to Tenover *et al.* J. Clin. Endocrinol and Metab. 65:1118 (1987).

[0048] Patients are given daily dosages of 1.5 mg/kg clomiphene; wherein the ratio of *trans*-clomiphene to *cis*-clomiphene is greater than 1. Patients are monitored for testosterone levels such that the dosage amount and dosage frequency may be adjusted to achieve therapeutic levels of testosterone in the patient.

Claims:

1. Use of a composition comprising 0% to 29% w/w of *cis*-clomiphene and 100% to 71% trans-clomiphene or pharmaceutically acceptable salts or solvates thereof and optionally one or more pharmaceutically acceptable diluents, adjuvants, carriers or excipients, for the manufacture of a medicament for treatment of a human male to increase serum levels of testosterone in the male.
2. Use according to claim 1 wherein the composition consists essentially of an effective amount of trans-clomiphene or a pharmaceutically effective salt or solvate thereof and optionally one or more pharmaceutically acceptable diluents, adjuvants, carriers or excipients.
3. Use according to claim 2, wherein the medicament is for administration at a dosage of between 1 to 200 mg of trans-clomiphene.
4. Use according to any one of claims 1 to 3, wherein the medicament is in the form of a filled capsule for oral use.
5. Use according to any one of claims 1 to 4, wherein the medicament is for treatment of erectile dysfunction, oligospermia, azoospermia, progressive decrease in muscle mass, fatigue, depressed mood or osteoporosis.

6. A filled capsule for oral use, comprising a composition comprising 0% to 29% w/w of cis-clomiphene and 100% to 71% trans-clomiphene or pharmaceutically acceptable salts or solvates thereof and optionally one or more pharmaceutically acceptable adjuvants, carriers or excipients.

FIG. 1

Normal Secretory Total Serum Testosterone Profiles in Healthy Young and Older Men

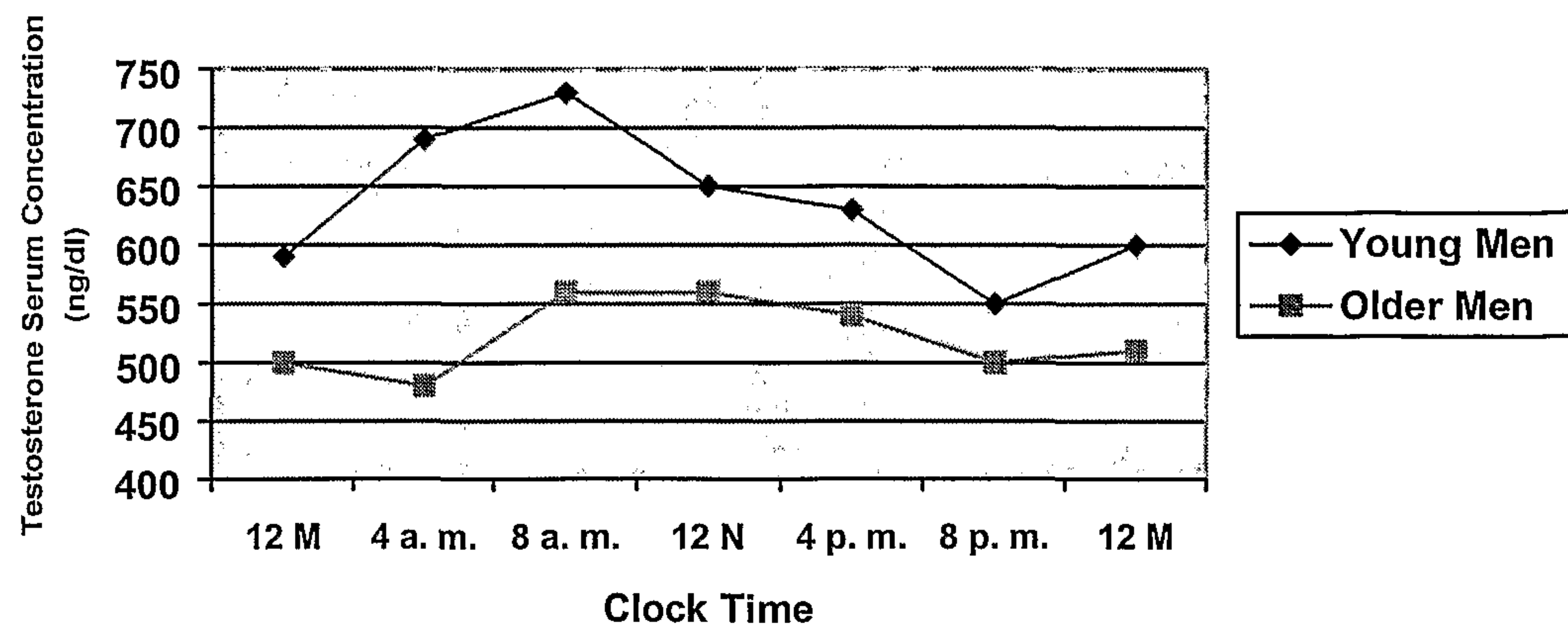


FIG. 2

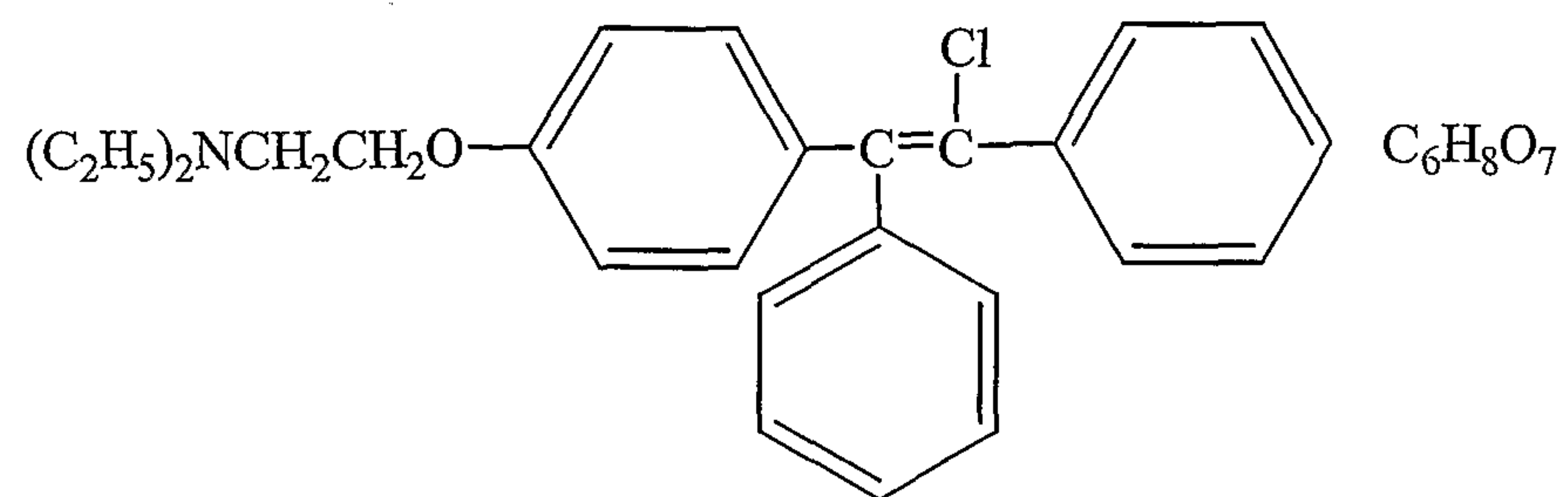


FIG. 3

