

9:45 - 10:30 am

Testosterone Replacement Therapy for Hypogonadism:
What is the Evidence? Is it Safe?

SPEAKER
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# primed

#### **Presenter Disclosure Information**

The following relationships exist related to this presentation:

Culley Carson, MD, FACS: Consultant for AbbVie Inc.; American Medical Systems; and Auxilium. Speaker for American Medical Systems and Auxilium.

#### Off-Label/Investigational Discussion

► In accordance with pmiCME policy, faculty have been asked to disclose discussion of unlabeled or unapproved use(s) of drugs or devices during the course of their presentations.

## **Learning Objectives**

- · Review role of testosterone in Men's Health
- Discuss the diagnosis of Hypogonadism
- Highlight published treatment guidelines for testosterone replacement therapy (TRT)
- Discuss recent data on cardiovascular risk factors for TRT
- Discuss the effects of TRT on the prostate in the aging male

## **Case Study**

- 59-year-old man, retired taxi driver presents for periodic follow up of hypertension
- · Type 2 diabetes for 10 yrs
- Fasting glucose between 108 mg/dl and 142 mg/dl
- · Hypertension
- · Hyperlipidemia
- · Erectile dysfunction for 2 years
- Low sexual desire

## Case Study - contd.

#### Symptoms

- Frequent and easy fatigue
- Decreased exercise
- Difficulty concentrating
- Nocturia
- Snoring

## Case Study - contd.

#### Medications

- Glipizide 5 mg bid
- Metformin 1000 mg bid
- Atorvastatin 40 mg before bedtime
- Benazepril 40 mg daily
- Aspirin 81 mg daily
- Tadalafil 20 mg prn

### Case Study - contd.

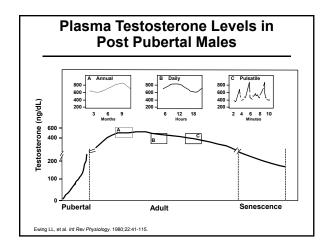
- · Physical examination
  - Height 69 inches, weight 215 lbs (BMI 33)
  - Waist circumference 42 inches
  - Normal hair distribution
  - Genital examination normal, including normal size testicles
  - Digital rectal exam: moderately enlarged prostate, no nodules
  - No other pathologic signs

# Case Study - contd.

- Labs
  - HbA1C 8.2%
  - Total cholesterol 143 mg/dl
  - LDL cholesterol 93 mg/dl
  - Normal blood urea nitrogen and creatinine
  - Microalbumin/creatinine in urine <10
  - Total serum testosterone 176 ng/dl

## Male Hypogonadism

- Endocrine disorder with:
  - Decreased testosterone and/or sperm production
  - Signs and symptoms of androgen deficiency
- · Causes of hypogonadism
  - Primary: Testicular defects
  - Secondary: Hypothalamic-pituitary defects
  - Combined (e.g, age-related decline, sickle cell disease)



# **Classical Hypogonadism**

- Congenital or acquired causes that may lead to irreversible testosterone deficiencies
  - Primary Hypogonadism
    - e.g., Klinefelter Syndrome, testicular trauma, cryptorchidism
  - Secondary Hypogonadism
    - e.g. Kallman Syndrome, pituitary tumor, prolactinomas

# **Biochemical Definitions of** Hypogonadism

#### Biochemical Definitions of Hypogonadism\*

Guidelines	nmol/l	ng/ml	ng/dl
EAA, ISA, ISSAM, EAU, ASA	Mild <12 Severe <8	<3.40 <2.31	<340 <231
Endocrine Society	<10.4	<3.00	<300
AACE	<7	<2.00	<200

\*According to international societies.

AACE = American Association of Clinical Endocrinologists; ASA = American Society of Andrology; EAA = European Acade
of Andrology; EAU = European Association of Utrology; ISA = International Society of Andrology; ISSAM = International
Society for the Study of the Aging Male.
Corona C. & Maggin M. (2009). ARI Rev Uro! doi:10.1038/nrurol.2009.235.

# **Dr Phil's Simple Testosterone Test**

According to Dr. Phil, men with a ring finger longer than their index finger, is an indication of higher testosterone levels

# **Conditions Associated with Testosterone Deficiency**

#### Suggest Evaluation

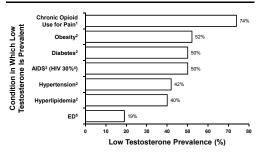
- · Sellar mass or irradiation with hypopituitarism
- · Conditions with medications (Glucocorticoids, Opioids)
- Cancer therapies chemotherapy, radiation
- HIV/cancer-associated weight loss
- · Osteoporosis/low trauma fracture (young men)

#### Suggest evaluation in presence of signs/symptoms

- · End stage renal disease and maintenance hemodialysis
- Type 2 diabetes mellitus
- · Moderate to severe COPD

Bhasin S, et al. J Clin Endocrinol Metab. 2010;95(6):2536-2559

# **Prevalence of Low Testosterone Other Conditions**



ED = erectile dysfunction; HIV = human immunostificiency virus.

1. Daniell HIV. | Phan. 2002;3(5):737-38.4. | Auliginan 1; et al. Int J Clin Pract. 2006;60(7):762-3. Grinspoon S, et al. Ann Intern Med. 1998;12(1):18-26. 4. Dobs AS. Ballieres Clin Endocrinol Metab. 1998;12(3):379-390. 5. Bodie J, et al. J Uric 2003;19(9):2282-2284.

# Signs and Symptoms of Hypogonadism

#### More Specific

- · Reduced or diminished
  - Sexual development
  - Libido
  - Spontaneous erections

  - Fertility - Secondary sexual
  - characteristics - Bone Mineral Density
- Increased
  - Fragility fractures
- Hot flushes, sweats

#### Less Specific

- · Reduced or diminished
  - Energy and vitality
  - Mood
  - Concentration and memory
  - Physical performance
  - Muscle bulk / strength
- Increased
  - Body fat

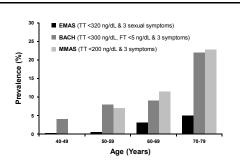
Bhasin S, et al. J Clin Endocrinol Metab. 2010;95(6):2536-2559.

# **Age-Related Decline of Testosterone**

- Testosterone decreases 1-2% per year after age 40
- · Institute of Medicine (2004): Insufficient data on TRT in elderly men
- Endocrine Society recommends caution in treating older men given paucity of data
- · Ongoing T-Trials to characterize benefits in elderly men

Araujo & Wittert. Best Pract Res Clin Endocrinol Metab. 2011;25(2):303-19.

# Prevalence of Symptomatic Hypogonadism

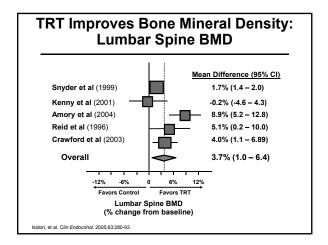


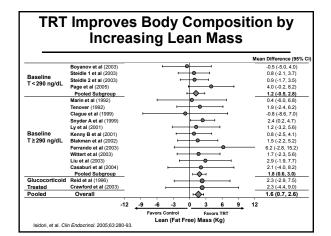
EMAS: Wu, et al. NEJM. 2010;363:123-135. BACH: Araujo, et al. J Clin Endocrinol Metab. 2007;92(11):4241-7. MMAS: Araujo, et al. J Clin Endocrinol Metab. 2004;89(12):5920-6.

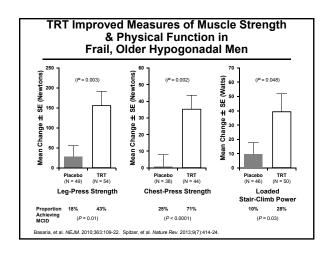
# TRT Has Potential Benefits in Multiple Body Systems

#### **Potential Benefits**

- · Bone Mineral Density
- · Lean Mass / Fat Mass
- · Muscle Strength and Physical Function
- · Sexual Function
- Mood
- · Fatigue





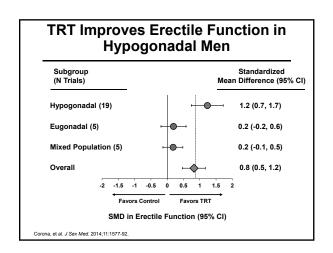


# Hypogonadism and ED Intrinsically Tied

- Hypogonadism common in men with ED1
  - Shared pathophysiology and comorbidities
- Clinicians need to screen and treat simultaneously<sup>1,2</sup>
- Testosterone combined with PDE5 inhibitors  $\underline{\textbf{may}}$  improve treatment outcomes  $^2$
- Chronic PDE5 inhibition improves endothelial function with sustained effect<sup>3</sup>
  - Rationale for vascular rehabilitation and cardioprotection

ED = erectile dysfunction; PDE5 = phosphodiesterase type 5.

1. Köhler 1S, et al. Urology. 2008;71(4):893-897. 2. Corona G, Maggi M. Nat Rev Urol. 2010;7(1):46-56. 3. Aversa A, et al. Irrl J Impor Res. 2007;19(2):200-207.



## **TRT: Mood and Fatigue**

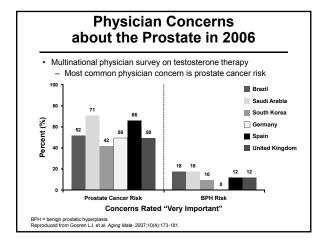
#### Mood

- Meta-analysis of 16 randomized trials (N=944) of heterogeneous population found moderate overall effect on mood
- · Effect significant in hypogonadal, but not in eugonadal subjects
- Effect significant in studies with mean age <60, but not ≥60

#### Fatique:

- Results vary across studies and patient populations
- Observational study of 799 men treated with testosterone showed a 22% reduction in fatigue scores over 6 months
- In a placebo-controlled study of HIV-infected men, a similar reduction in fatique scores was observed over 8 weeks

Amanatkar & Chibnall. Ann Clin Psychiatry. 2014;26(1):19-32. Rabkin, et al. J Clin Psychopharm 1999. Pexman-Fieth, et al. Aging Male. 2014.



#### **Testosterone and the Prostate**

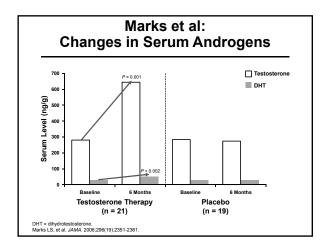
- Testosterone therapy significantly affects PSA concentrations at low levels of serum testosterone, it does <u>not</u> appear to affect prostate size or prostatic testosterone levels
  - Perhaps due to early saturation of androgen receptors within prostate
- <u>No</u> evidence that testosterone therapy causes new prostate cancer in hypogonadal men
- To date, 283 men reported in literature have received testosterone therapy after prostate cancer treatment, with low recurrence rates of 1.4%

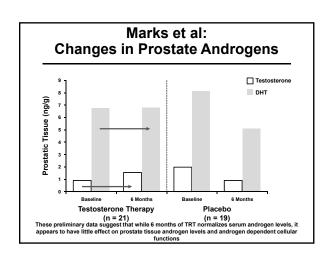
PSA = prostate-specific antigen

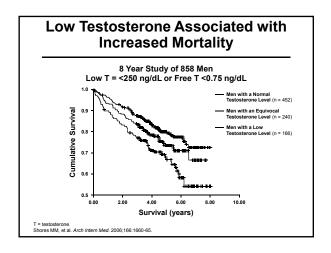
# Prostate Cancer and Testosterone Therapy FACTS

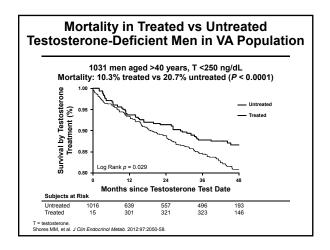
- Fear of causing prostate cancer leaves many appropriate patients untreated
- No evidence of causality of testosterone use and development of prostate cancer
- Testosterone will stimulate growth of existing prostate cancers
- · Obtain consult for any concern
  - PSA abnormal per guidelines
  - Abnormal prostate exam

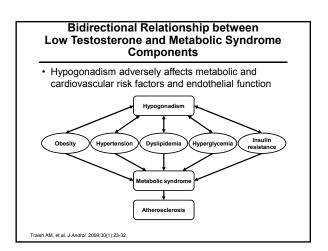
PSA = prostate-specific antigen.
1. Gooren I.J, et al. *Aging Male*. 2007;10(4):173-181. 2. Rhoden EL, et al. *N Engl J Med*. 2004;350(5):482-492. 3. Raynaux *JP*. *J Sterodi Biothem Mol Biol*. 2006;102(1-5):261-266. 4. Wang C, et al. *J Androl*. 2009;30(1): 1-9. 5. Carroll P, et al. *Urology*. 2001;57(2):217-224.

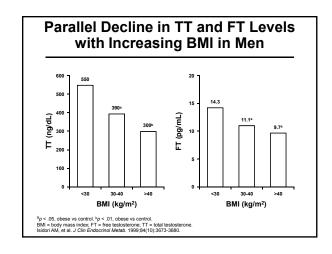


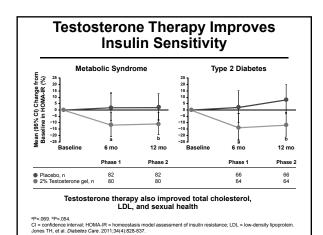












# Mixed Epidemiologic Results in Studies of TRT and Mortality among Men with Low Testosterone

- Vigen et al. (2013)
  - -N = 8,709 with testosterone <300 ng/dL
  - Endpoint: all-cause mortality, MI, or stroke
  - Adjusted HR = 1.29 (95% CI: 1.05, 1.58)
- Shores et al. (2012)
  - N = 1,031 with testosterone ≤250 ng/dL
  - Endpoint: all-cause mortality
  - Adjusted HR = 0.61 (95% CI: 0.42, 0.88)

Vigen, et al. JAMA. 2013;310(17):1829-36. Shores, et al. J Clin Endocrinol Metab. 2012;97(6):2050-8.

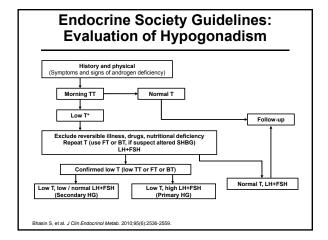
# Mixed Epidemiologic Results in Studies of TRT and MI among Men with Low Testosterone

- Finkle et al. (2014)
  - N = 55,593 men on TRT examining non-fatal MI
  - Men ≥65: RR = 2.2 (95% CI: 1.3, 3.8)
  - Men <65 with history of HD: RR = 2.9 (95% CI: 1.5, 5.6)
  - Remaining 80% of cohort: RR = 0.9 (95% CI: 0.6, 1.3)
- Baillargeon et al. (2014)
  - N = 6,355 TRT matched to 19,065 non-TRT controls
  - Men >65: HR = 0.84 (0.69, 1.02)
  - TRT more protective against MI in men at higher MI risk

Finkle, et al. PLoS ONE. 2014;9(1):e85805. Baillargeon, et al. Ann Pharmacother. 2014;48(9):1138-44.

# Recent Epidemiological Studies on TRT and Mortality/CV Outcomes

- · Limitations
  - Heterogeneity of study populations, intervention duration, and study designs
  - Variable definitions, ascertainment of CV outcomes
  - Unclear treatment indications, treatment regimens, testosterone levels, and exposure
  - Residual confounding
  - Study groups differed in baseline CV risk factors
- · Overall Assessment
  - Inconsistent findings across studies
  - Difficult to draw inferences



# Testosterone Formulations and Regimens Testosterone Formulation Injectable Testosterone Formulation Injectable Testosterone Formulation Testosterone Formulation Delatestryl, Depo-Testosterone Depo-Testosterone Testosterone undecancatel\* Testosterone Testos

# Endocrine Society Guidelines: Monitoring of TRT

	Baseline	Each Visit	3-6 Months	Annually	1-2 Years
Symptom response		X	X	X	
Adverse events		Х	Х	Х	
Testosterone levels	Х		Х		
Hematocrit	Х		Xa	Х	
BMD of lumbar spine / femoral neck					Xp
Digital rectal exam/PSA	Х		Xc		

"If hematocrit >54%, stop therapy until hematocrit decreases to safe level; evaluate patient for hypoxia and OSA, reinitiat therapy at lower dose. "For patients with osteporosis or low trauma tracture, consistent with regional standard of care. "After 34 months, perform in accordance with guidelines for prostate cancer screening, depending on age and race of patients. Obtain unological consultation under certain conditions.

Bhasin S, et al. 20 int Endocrinut Media. 2010;45(9):258-2559.

# **Summary of 2010 Endocrine Guidelines**

Do Not Treat

- Patients with breast or prostate cancer
- A palpable prostate nodule or induration
- Abnormal PSA
- · Consider consultation in high risk patients
- Patients with erythrocytosis
- Untreated severe sleep apnea
- Severe lower urinary tract symptoms with International Prostate Symptom Score > 19
- Uncontrolled or poorly controlled heart failure

Bhasin S, Cunningham GR, Hayes FJ, et al. J Clin Endocrinol Metab . 2010, 96(6): 2536-2559

# Conclusions: Benefits Determined by Patient Selection

- TRT is well accepted in men with classical signs/symptoms & etiologies of hypogonadism
- Select chronic diseases & medications result in suppression of the H-P-G axis (e.g. opiates, radiation therapy) & may lead to symptoms & consideration of treatment
- Poorly defined, mild reductions from chronic disease & aging should be a risk-benefit discussion with frequent re-evaluations