Studies in JSM you must know about

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Editor-in-Chief

The Journal of Sexual Medicine

Relevant Rigorous Reproducible Readable

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 - 45 areas of sexual medicine
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State of the Journal

- Submissions (840 in 2015)
- Impact factor (3.15)
- Ranked 17/76 in U+N journals
- Ranked 1st among sexual medicine journals
- 67% rejection rate (19% submitted to OA journal)
- Turnaround time = 21 days
- YOUR JOURNAL

Electronic Journal Capabilities

- Interactivity enhancing the journal experience
- Downloadable content
- Referencing searching within manuscript
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Endorsement of Models Describing Sexual Response of Men and Women with a Sexual Partner: An Online Survey in a Population Sample of Danish Adults Ages 20–65 Years

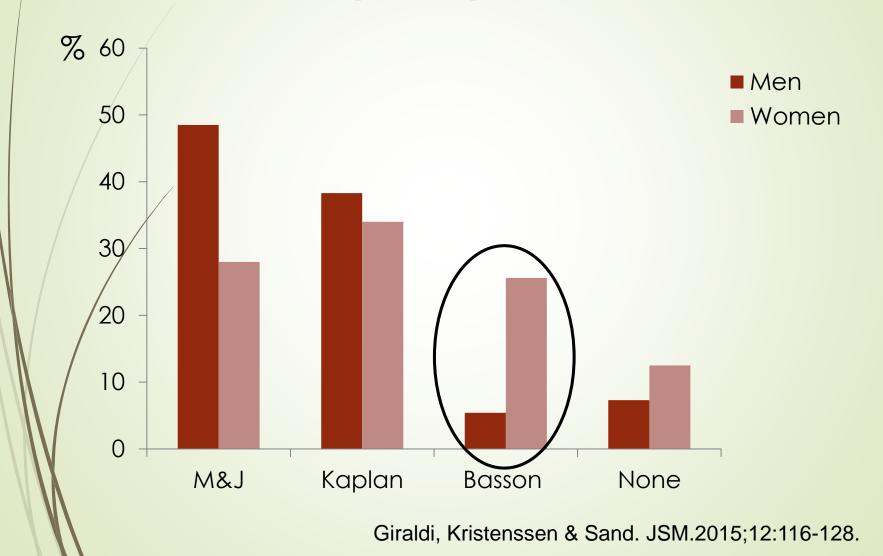
- Annamaria Giraldi, MD, PhD, Ellids Kristensen, MD, and Michael Sand, PhD
- J Sex Med. 2015;12:116-28.

Which model is best?

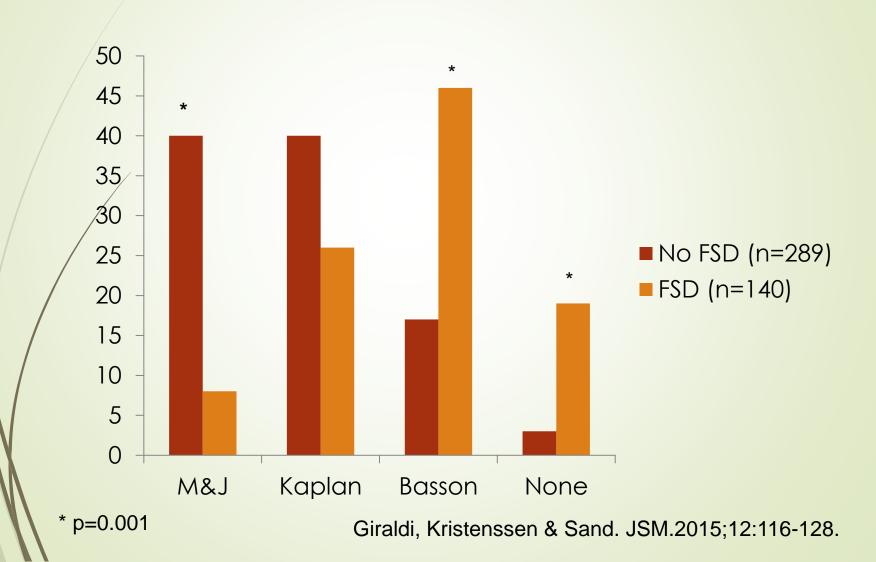
Methods

- 1996 Danish women aged 20-65 years mailed
 84-item questionnaire, 573 responded (29%)
- ■2004 Danish men aged 20-65 years mailed a 91item questionnaire,499 responded (25%)
- Parameters assessed
 - Demographic data
 - Perceptions of models of sexual response
 - Sexual function (FSFI ♀), (IIEF-15 domain ♂)
 - Sexual distress (FSDS ♀)
 - Satisfaction with sexual life

Models endorsed by men (n=410) & women (n=465)



Effect of Female Sexual Dysfunction (FSD) (FSFI <26.55)



Conclusions

- Majority of men and women endorse the M&J or Kaplan model.
- Choice of the M&J is correlated to being a man, Basson to being a woman
- In both men and women with sexual dysfunction, significantly more choose the Basson model or none of the models, but men still prefer the other models.
- There is no difference in women with/without dysfunction chosing the Kaplan model
- Basson model is correlated to being dissatisfied with sexual life and for women with living together with partner.
- No correlation to length of relationship, sexual abuse, depression, smoking, BMI and frequency of sexual activity

No model fits all

Clinical implications

- Focus on different types of desire receptive desire is also desire
- Focus on differet models within the couple
- Sex therapy focus on intimacy rather than genital response to prevent performance anxiety.
- May suit women better than men, who still focus on the genital response

ORIGINAL RESEARCH—ENDOCRINOLOGY

Is There a Correlation Between Androgens and Sexual Desire in Women?

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DOI: 10.1111/jsm.12774

J Sex Med 2015;12:358-373

Methods

- Cross-sectional investigation of 560 healthy women, aged 19-65 years
- Female Sexual Function Index (FSFI)
 - ■Total score of desire domain
- Hormonal measurements of:
 - Total and free testosterone
 - Androstendione
 - Dehydroepiandrosteronsulfate (DHEAS)
 - Primary androgen metabolite: androsterone glucuronide (ADT-G).
 - Androstendione / total testosterone*

^{*} Estimated enzymantic activity of 17b-hydroxysteriod dehydrogenase

Correlation between hormonal level and sexual desire, adjusted for age

	Sexual desire Total cohort (n = 560)	Sexual desire No use of HC/HRT* (n = 346)
Total testosterone, nmol/L	ns	ns
Free testosterone, nmol/L	p < 0.05	p < 0.05
Androstendione, nmol/L	p < 0.01	p < 0.01
DHEAS, nmol/L	ns	p < 0.01
ADT-G, nmol/L	ns	ns
Androstendione/ total testosterone ratio	ns	p < 0.05

Age Stratification Results

- Both total and free testosterone, androstenedione and DHEAS statistically significantly correlated with sexual desire in women aged 25–44 years with no use of systemic HC.
- In women aged 45-65 years, androstenedione (p=0.001) and androstenedione:total testosterone ratio (p<0.001)positively correlated with sexual desire</p>
- ADT-G was not correlated with sexual desire in any of three age groups.
- Sexual desire declined with age (p< 0.001, p=0,024) in both pre- and postmenopausal women.

Conclusion

- Biosynthesis of testosterone is decreased in women who experience low desire.
- First study to show correlation androgen precurors and desire
- Speed of transformation of androstendione to testosterone may influence older womens sexuality

Clinical Implications

Androgens are important to women's sexual desire, especially for women aged 25–44 years.

Possible relevance to measurement androgens (total and free testosterone, androstenedione, and DHEAS) using mass spectrometry in women complaining about a lack of sexual desire in this age group

Psychological Treatment Trials for Hypoactive Sexual Desire *Disorder*

- Robert E. Pyke, MD, PhD; Anita Clayton, MD
- J Sex Med 2015;12:2451-2458.

Study Aim and Methods

- Aim: evaluate published controlled trials of CBT and MMT (Mindfulness Meditation Training) for sexual desire disorders
- Perspective: sexual medicine standards of control paradigms, risk/benefit ratios, clinical significance
- Medline 10 years; evaluate study quality via 10 metrics and efficacy as mean change, proportion of responders and remitters

Benefit-Risk Issues

- Behavioral measure should ground improvements in HSDD to the patient's sexual life
- Benefit should measure, on well-validated outcome endpoints, sexual desire and distress, sexual activity
- Validated instruments for measuring harms of HSDD such as marital discord and failing relationships, are not available.
- Thus treatments for HSDD should have minimal or no side effects to show positive risk-benefit ratio.
- Favors psychological therapies however, controlled research has not supported any form of psychological therapy

Results

- Three controlled trials support CBT
- Two controlled trials support MMT
- Reports of trials each lacked several scientific requirements:
 - Hierarchy of endpoints with planned primary endpoint
 - Sufficient information on endpoint to reproduce it
 - Randomization
 - "Open label" nature and lack of blinded rates
 - Adequate control waitlist?
 - Accepted measures of benefits and harms
 - Compliance data
 - Outcomes of clinical relevance: meaningful benefit, clinical significance, proportion of responders and remitters on well-validated measures

Conclusions and Implications

- Psychological treatments for HSDD are not yet supported by adequate clinical trials.
- Current scientific and regulatory standards for drug trials should be applicable to psychological treatment.