

# Diagnostic evaluation of the infertile male: a committee opinion

Practice Committee of the American Society for Reproductive Medicine

American Society for Reproductive Medicine, Birmingham, Alabama

The purpose of this ASRM Practice Committee report is to provide clinicians with principles and strategies for the evaluation of couples with male infertility problems. This revised document replaces the document of the same name, last published in 2012 (Fertil Steril 2012;98:294–301). (Fertil Steril® 2015;103:e18–e25. ©2015 by American Society for Reproductive Medicine.)

Earn online CME credit related to this document at [www.asrm.org/elearn](http://www.asrm.org/elearn)

**Discuss:** You can discuss this article with its authors and with other ASRM members at <http://fertilityforum.com/asrmpraccom-diagnostic-evaluation-infertile-male/>



Use your smartphone to scan this QR code and connect to the discussion forum for this article now.\*

\* Download a free QR code scanner by searching for "QR scanner" in your smartphone's app store or app marketplace.

Approximately 8%–15% of couples are unable to conceive after 1 year of unprotected intercourse (1). A male factor is solely responsible in ~20% of infertile couples and contributes in another 30%–40% of couples (2). A male infertility factor is often defined by abnormal semen parameters but may be present even when the semen analysis is normal. The purpose of this document is to provide clinicians with principles and strategies for the evaluation of couples with male infertility problems.

## GOALS OF EVALUATION

Male infertility can be due to a variety of conditions, many, but not all, of which can be identified and treated. When the cause of abnormal semen parameters cannot be identified, as is true in many patients, the condition is termed idiopathic. Rarely, patients with normal semen quality may have sperm that either are incapable of oocyte fertilization or harbor genetic abnormalities that prevent normal fetal development.

Ideally, the identification and treatment of correctable conditions

will improve the male partner's fertility and allow conception to be achieved naturally. Detection of certain genetic causes of male infertility provides the opportunity to inform affected couples about the risk of transmitting genetic abnormalities that may affect the health of offspring, may affect the chance for successful treatment, and can help to guide treatment options. Evaluation of the infertile man also is aimed at identifying any underlying medical conditions that may present as infertility. Some, such as testicular cancer and pituitary tumors, can have serious health consequences if not properly diagnosed and treated (3).

## INDICATIONS FOR EVALUATION

Evaluation for infertility is indicated for couples who fail to achieve a successful pregnancy after  $\geq 12$  months of regular unprotected intercourse. Earlier evaluation and treatment may be justified, based on medical history and physical findings and is warranted after 6 months for couples in which the female partner is  $>35$  years old (4). Men

having concerns about their future fertility also merit evaluation.

At a minimum, the initial screening evaluation of the male partner of an infertile couple should include a reproductive history and analysis of at least one semen sample. If the initial evaluation is abnormal, then referral to someone experienced in male reproduction is recommended.

## Reproductive History

The reproductive history should include: 1) coital frequency and timing; 2) duration of infertility and previous fertility; 3) childhood illnesses and developmental history; 4) systemic medical illnesses (such as diabetes mellitus and upper respiratory diseases); 5) previous surgery; 6) medications and allergies; 7) sexual history (including sexually transmitted infections); and 8) exposures to gonadotoxins (including environmental and chemical toxins and heat). Previous fertility does not exclude the possibility of a newly acquired, secondary, male infertility factor. Evaluation is the same for men with primary infertility (never having fathered a pregnancy) and secondary infertility (having previously fathered a pregnancy).

## Semen Analysis

Semen analysis is the cornerstone of the laboratory evaluation of the

Received December 4, 2014; accepted December 5, 2014; published online January 15, 2015.

Reprint requests: Practice Committee, American Society for Reproductive Medicine, 1209 Montgomery Hwy., Birmingham, Alabama 35216 (E-mail: [ASRM@asrm.org](mailto:ASRM@asrm.org)).

Fertility and Sterility® Vol. 103, No. 3, March 2015 0015-0282/\$36.00

Copyright ©2015 American Society for Reproductive Medicine, Published by Elsevier Inc.

<http://dx.doi.org/10.1016/j.fertnstert.2014.12.103>

This page was deleted because of demo limitations.

47. Oates RD, Silber S, Brown LG, Page DC. Clinical characterization of 42 oligospermic or azospermic men with microdeletion of the AZFc region of the Y chromosome, and of 18 children conceived via ICSI. *Hum Reprod* 2002;17:2813–24.
48. Brandell RA, Mielnik A, Liotta D, Ye Z, Veeck LL, Palermo GD, et al. AZFb deletions predict the absence of spermatozoa with testicular sperm extraction: preliminary report of a prognostic genetic test. *Hum Reprod* 1998;13:2812–5.
49. Krausz C, Quintana-Murci L, McElreavey K. Prognostic value of Y deletion analysis: what is the clinical prognostic value of Y chromosome microdeletion analysis? *Hum Reprod* 2000;15:1431–4.
50. Kent-First MG, Kol S, Muallem A, Ofir R, Manor D, Blazer S, et al. The incidence and possible relevance of Y-linked microdeletions in babies born after intracytoplasmic sperm injection and their infertile fathers. *Mol Hum Reprod* 1996;2:943–50.
51. Jorgez CJ, Weedin JW, Sahin A, Tannour-Louet M, Han S, Bournat JC, et al. Aberrations in pseudoautosomal regions (PARs) found in infertile men with Y-chromosome microdeletions. *J Clin Endocrinol Metab* 2011;96:E674–9.
52. Kent-First M, Muallem A, Shultz J, Pryor J, Roberts K, Nolten W, et al. Defining regions of the Y-chromosome responsible for male infertility and identification of a fourth AZF region (AZFd) by Y-chromosome microdeletion detection. *Mol Reprod Dev* 1999;53:27–41.
53. Carrell DT. The clinical implementation of sperm chromosome aneuploidy testing: pitfalls and promises. *J Androl* 2008;29:124–33.
54. Egozcue S, Blanco J, Vendrell JM, Garcia F, Veiga A, Aran B, et al. Human male infertility: chromosome anomalies, meiotic disorders, abnormal spermatozoa and recurrent abortion. *Hum Reprod Update* 2000;6:93–105.
55. Carrell DT, Wilcox AL, Lowy L, Peterson CM, Jones KP, Erickson L, et al. Elevated sperm chromosome aneuploidy and apoptosis in patients with unexplained recurrent pregnancy loss. *Obstet Gynecol* 2003;101:1229–35.
56. Petit FM, Frydman N, Benkhalifa M, Le Du A, Aboura A, Fanchin R, et al. Could sperm aneuploidy rate determination be used as a predictive test before intracytoplasmic sperm injection? *J Androl* 2005;26:235–41.
57. Tempest HG, Martin RH. Cytogenetic risks in chromosomally normal infertile men. *Curr Opin Obstet Gynecol* 2009;21:223–7.