

# The management of obstructive azoospermia: a committee opinion

Practice Committee of the American Society for Reproductive Medicine in collaboration with the Society for Male Reproduction and Urology

American Society for Reproductive Medicine, Birmingham, Alabama

Infertility due to obstructive azoospermia may be treated effectively by surgical reconstruction or by retrieval of sperm from the epididymis or testis, followed by in vitro fertilization with intracytoplasmic sperm injection. This replaces the ASRM documents titled “Sperm retrieval for obstructive azoospermia” and “The management of infertility due to obstructive azoospermia,” last published in 2008. (Fertil Steril® 2019;111:873–80. ©2019 by American Society for Reproductive Medicine.)

**Discuss:** You can discuss this article with its authors and other readers at <https://www.fertstertdialog.com/users/16110-fertility-and-sterility/posts/43953-27784>

## OBSTRUCTIVE AZOOSPERMIA

Obstructive azoospermia is the result of a blockage of the male reproductive tract, leading to a complete absence of sperm in the ejaculate, and accounts for approximately 40% of all cases of azoospermia (1). Obstruction may be congenital or acquired and may include one or more segments of the male reproductive tract: epididymis, vas deferens, and ejaculatory ducts. Congenital causes of obstructive azoospermia include congenital bilateral absence of the vas deferens (CBAVD) and idiopathic epididymal obstruction. Acquired causes of obstructive azoospermia include vasectomy, infection, trauma, or iatrogenic injury. The evaluation of nonobstructive azoospermia is covered in another document (2).

## MICROSURGICAL RECONSTRUCTION

Microsurgical techniques for the treatment of obstructive azoospermia were first introduced by Silber and Owen in 1977 and are considered the gold standard for reconstructive surgery involving the male reproductive tract

(3–5). Scrotal or inguinal vasovasostomy may be employed for vasal obstruction secondary to vasectomy, iatrogenic vasal injury due to inguinal or scrotal surgery such as herniorrhaphy or hydrocelectomy, or solitary vasal obstruction secondary to infection or trauma. Vasovasostomy is not feasible in the setting of multifocal obstruction along the vas deferens.

Vasoepididymostomy is indicated for idiopathic epididymal obstruction, secondary epididymal obstruction due to long-standing vasal obstruction, or iatrogenic epididymal obstruction following interventions such as epididymal aspiration. Naturally, multifocal epididymal obstruction necessitates that the level of the anastomosis be proximal to all sites of obstruction.

The primary indication for vasovasostomy or vasoepididymostomy is to restore fertility in the setting of obstructive azoospermia. Although the use of these procedures has been described for the treatment of postvasectomy pain syndrome (6), affected men should be offered counseling or conservative methods of pain management prior to consideration of these reconstructive procedures.

## SURGICAL TECHNIQUE

### Incision

Scrotal vasovasostomy and vasoepididymostomy are usually performed through 2–3 cm vertical incisions in the anterior aspect of the hemiscrotum. When greater exposure or mobilization of the vas is required, for example, following a high vasectomy or loss of a large segment of the scrotal vas deferens, the scrotal incisions can be extended into the inguinal region. Alternatively, the surgeon can use an infrapubic incision to better mobilize the vas deferens (7) to perform a tension-free anastomosis.

Inguinal vasovasostomy may be performed via an inguinal approach if the vasal obstruction is focal and easily identified in the inguinal canal, as for example, following iatrogenic injury to the vas during inguinal surgery. Alternatively, laparoscopic or robotic assistance may be required for identification and dissection of the abdominal vas deferens prior to performing a microsurgical anastomosis.

### Intraoperative Considerations

For men undergoing vasectomy reversal, the decision to perform either a vasovasostomy or a vasoepididymostomy is made intraoperatively. As such, vasectomy reversal should ideally be

Received February 6, 2019; accepted February 12, 2019.

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

Fertility and Sterility® Vol. 111, No. 5, May 2019 0015-0282/\$36.00

Copyright ©2019 American Society for Reproductive Medicine, Published by Elsevier Inc.

<https://doi.org/10.1016/j.fertnstert.2019.02.013>

undertaken by surgeons who are skilled in both vasovasostomy and vasoepididymostomy. Although obstructive interval (8), length of the testicular vasal remnant (9), sperm granuloma (10), vasectomy site (11), prior vasectomy reversal, and surgeon skill are all factors to consider when making this decision, the quality of the vasal fluid remains the single most important determinant of whether vasovasostomy or a vasoepididymostomy should be performed. Testicular vasal fluid may be watery and copious or thick and creamy in consistency and have one of the following microscopic characteristics:

Grade 1 – mainly normal motile sperm

Grade 2 – mainly normal non-motile sperm

Grade 3 – mainly sperm heads

Grade 4 – only sperm heads

Grade 5 – no sperm

Vasovasostomy should be performed for grades-1–4 vasal fluid (12, 13). When no sperm are observed in the vasal fluid, the likelihood of return of sperm to the ejaculate is greatest when the fluid is watery and copious, and lowest when it is thick and creamy (12). Therefore, for grade-5 vasal fluid, vasovasostomy should be performed if the fluid is watery and copious. If the fluid is thick and creamy, careful inspection of the epididymis under magnification can reveal a discolored or indurated area in the epididymis, signifying tubule rupture due to back pressure, or a demarcation between collapsed and dilated tubules. Vasoepididymostomy should be performed in both these circumstances.

Prior to anastomosis, patency of the distal length of the vas deferens should be confirmed by performing a saline vasogram, using a 24-gauge blunt-tipped peripheral venous catheter inserted directly into the lumen. Formal vasography is rarely necessary. Regardless of the location of the anastomosis, adequate mobilization of the vas, without devascularization, should be performed to ensure a tension-free anastomosis.

### Anastomotic Methods

Although some surgeons have described satisfactory results using macrosurgical techniques or loupe-magnification (14), it is generally accepted that microsurgical anastomoses yield better outcomes than macrosurgical anastomoses, particularly for vasoepididymostomy.

Most surgeons perform vasovasostomy using a two-layer microsurgical anastomosis, by first placing 5–8 interrupted 10-0 nylon sutures in the inner mucosal edges of the vas, incorporating a small portion of the inner muscular layer, followed by 5–8 additional interrupted 9-0 nylon sutures in the outer muscular layer (3). A modified one-layer anastomosis using 4–6 interrupted 9-0 nylon sutures through the full thickness of the vas has also been described (15, 16). A meta-analysis showed no statistically significant difference in outcomes between the techniques (16).

Several different techniques have been described for microsurgical vasoepididymostomy, including an end-to-side anastomosis (17), triangulation (18), tubular invagi-

nation (19), and tubular intussusception techniques (20). In all techniques, the epididymal tubule is pulled up into the lumen of the vas deferens. The intussuscepted end-to-side vasoepididymostomy is currently the most commonly employed technique.

Robotic-assisted surgery has been applied for scrotal vasovasostomy and vasoepididymostomy as an alternative to microsurgery, but its use is not widespread at present (21). Robotic-assisted intra-abdominal vasovasostomy, using the formal two-layer anastomosis discussed above, has also been described as a novel approach for repairing intra-abdominal vasal defects that minimizes morbidity and obviates the need for an operating microscope (22, 23).

### Sperm Retrieval and Cryopreservation

Patients should be counseled about the option of intraoperative sperm retrieval for cryopreservation via testicular biopsy or aspiration at the site of a vaso-vasal or vaso-epididymal anastomosis if microsurgical reconstruction is unsuccessful. This option may be more relevant for patients with a high preoperative likelihood of requiring a bilateral vasoepididymostomy. Costs of cryopreservation should also be discussed preoperatively.

### POSTOPERATIVE CARE

Most surgeons will advise patients to use ice packs in the immediate postoperative period to minimize pain and scrotal swelling. A scrotal supporter should continue to be used for 3–4 weeks after surgery. Heavy physical activity, including weight lifting, straddle-type activities, sexual intercourse, and ejaculation, should also be avoided for 3–4 weeks after surgery. Pain is usually well controlled with oral analgesics, both nonsteroidal anti-inflammatory drugs (NSAIDs) and narcotics.

### COMPLICATIONS

Complications after microsurgical reconstructive procedures are rarely reported in the published literature. Expected complications of scrotal surgery, such as scrotal hematomas, can usually be managed expectantly.

### MONITORING

Sperm may return to the ejaculate as early as 1 month after surgery, especially in the case of vasovasostomy. Semen analyses may be obtained every 8–12 weeks in the postoperative period until sperm concentration and motility return to normal or until a pregnancy occurs. Even after sperm parameters return to normal, semen analysis may be performed at least every 12 weeks until pregnancy occurs to promptly identify patients who may again become obstructed due to anastomotic failure or scarring.

The incidence of postoperative re-obstruction ranges from 12% (12/98) after vasovasostomy to approximately 21% (11/52) after vasoepididymostomy (24). Surgery is considered to have failed if sperm do not return to the ejaculate by 6 months after vasovasostomy or by 18 months after vasoepididymostomy (25). Most pregnancies occur within

24 months after surgery without any further intervention (12). If pregnancy fails to occur, despite normal semen quality and the lack of any identifiable female infertility factors, evaluation for antisperm antibodies may be considered.

## OUTCOMES

The Vasovasostomy Study Group reported a 97% patency rate (86/89) and 76% pregnancy rate (56/74) following vasovasostomy among patients who were less than 3 years out from vasectomy (12). In their report, these outcomes declined as the obstructive interval increased to a patency rate of 71% (32/45) and a pregnancy rate of 30% (11/37) at greater than 15 years (12). Although other studies have not shown a direct correlation between obstructive interval and patency rates after vasovasostomy, a significant decline in pregnancy rates has been noted for intervals greater than 15 years (26).

The Vasovasostomy Study Group also reported surgical success rates based on the quality of the vasa fluid. Patency and pregnancy rates ranged from 94% (116/123) and 63% (62/98), respectively, for quality of grade-1 fluid to 60% (50/83) and 31% (20/65), respectively, for quality of grade-5 fluid (12).

Techniques for vasoepididymostomy have evolved over time, with an evolution of surgical success rates as well. Patency rates for the currently used intussuscepted end-to-side vasoepididymostomy range from 80% (12/15) (20) to 84% (53/63) (27), with pregnancy rates of 40% (14/35) (27) to 44% (4/9) (20).

## MANAGEMENT OF OPERATIVE FAILURES

Men who remain azoospermic may be offered a repeat operation but should be counseled that repeat procedures are usually more technically challenging. According to findings from the Vasovasostomy Study Group, sperm returned to the ejaculate after repeat vasovasostomy in 75% of men (150 of 199 patients) and 43% of their partners subsequently achieved pregnancy (52 of 120 couples) (12). Other groups have reported comparable results (28–30).

After a failed vasoepididymostomy, a repeat procedure may not always be technically feasible. However, in the hands of experienced surgeons, acceptable success rates after repeat vasoepididymostomy are possible, with one series reporting an overall patency rate of 67% (12/18 men) and a natural conception rate of 25% (3/12 couples) (31).

## SPERM-RETRIEVAL TECHNIQUES

These techniques can be used in the setting of post-vasectomy obstruction, CBAVD, ejaculatory duct obstruction (EDO), and unreconstructable causes of obstructive azoospermia. These procedures do not yield sufficient sperm for intrauterine insemination (IUI) and rarely for assisted reproductive technology (ART), including in vitro fertilization (IVF) (32, 33). ART involves the manipulation of both ova and sperm outside the body and does not include IUI. Surgically retrieved testicular or epididymal sperm used for intracytoplasmic sperm injection (ICSI) have similar

outcomes whether fresh or frozen sperm are used: fertilization rates between 45% and 60% (8/17, 93/157) per injected oocyte; clinical pregnancy rates from 23%–35% (23/100, 5/17), and live-birth rates of approximately 18%–36% (14/77 and 7/19) (34–36). This is comparable to the use of ejaculated sperm (37).

Percutaneous epididymal sperm aspiration (PESA) is conducted in the office or laboratory procedure room under local anesthesia or monitored anesthesia care. It can be repeated and does not require microscopic guidance. After a spermatic cord block, the epididymis must be stabilized between the surgeon's forefinger and thumb. A butterfly needle with a 20 mL syringe is inserted into the caput epididymis. With gentle suction pressure, the needle is withdrawn until fluid enters the tubing. This is continued until sperm of adequate quality and quantity are obtained. If no sperm is aspirated, microsurgical epididymal sperm aspiration (MESA), testicular biopsy, or aspiration can still be performed.

## Microsurgical Epididymal Sperm Aspiration

Under general anesthesia or intravenous (IV) sedation, MESA techniques include individual epididymal incision and effluent aspiration or micropuncture of individual tubules (38, 39). Using an operating microscope, individual epididymal tubules are identified and then aspirated sequentially until optimal quantity and quality of sperm are obtained. Puncture sites are then closed or cauterized. Sperm of best quality are found near the testis in the proximal epididymis and are concentrated to about 1 million sperm/ $\mu$ L (40).

## Open Testicular Biopsy

Though the epididymis is a richer source of motile sperm than the testicle, testis retrieval can still be useful. There is no appreciable difference in pregnancy rate with ART between testicular or epididymal sperm for men with obstructive azoospermia (37, 41). Though sperm retrieval is similar, MESA may provide higher live-birth rates (42). Ideally, only one tunical incision is necessary to obtain sufficient tissue and sperm. Care must be taken to avoid damage to the testicular blood supply within the tunical wall.

## Percutaneous Testicular Sperm Aspiration (TESA)

After local cord block and stabilization of the testicle, a needle is inserted along the testicular long axis during TESA and then repeatedly redirected with gentle pressure and aspiration until tubules are sufficiently disrupted and adequate sperm are obtained. Fine-needle aspiration can also be used diagnostically and to procure sperm for ART.

## Percutaneous Testicular Biopsy (PercBiopsy)

A local spermatic cord block and a 14-gauge biopsy gun is used during a PercBiopsy to remove a small cylinder piece of testicular tissue. Multiple passes can be performed if needed.

TABLE 1

| Advantages and disadvantages of sperm-retrieval techniques. |                                                                                                                                         |                                                                                                                                    |
|-------------------------------------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------|------------------------------------------------------------------------------------------------------------------------------------|
| Technique                                                   | Advantages                                                                                                                              | Disadvantages                                                                                                                      |
| PESA                                                        | No microsurgical expertise required<br>Local anesthesia<br>Few instruments<br>Fast and repeatable<br>Minimal postoperative discomfort   | Few sperm retrieved<br>Risk of hematoma<br>Damage to adjacent tissue                                                               |
| MESA                                                        | Best clinical pregnancy rates<br>Large number of sperm retrieved<br>Excellent results with cryopreservation<br>Reduced risk of hematoma | Requires microsurgical expertise<br>Increased cost<br>General or local anesthesia<br>Incision required<br>Postoperative discomfort |
| TESE                                                        | No microsurgical expertise required<br>Local or general anesthesia<br>Few instruments<br>Fast and repeatable                            | Relatively few sperm retrieved<br>Limited risk of testicular atrophy (with multiple biopsies)                                      |
| PercBiopsy, TESA                                            | No microsurgical expertise required<br>Local anesthesia<br>Few instruments<br>Fast and repeatable<br>Minimal postoperative discomfort   | Few sperm retrieved<br>Risk of testicular atrophy<br>Risk of hematoma                                                              |

### Complications/Risks

Techniques for sperm retrieval are generally very safe, even under local anesthesia. The incidence of hematoma, persistent pain, swelling, or infection is low (43–45). The risk of associated birth defects after sperm retrieval with ICSI is similar to ICSI using ejaculated sperm (37).

### Cryopreservation

Sperm retrieved with these sperm-retrieval methods can be freshly used for IVF/ICSI. Alternatively, the sperm can be frozen and used later after thawing. Couples should be counseled that sperm can be lost in the freeze-thaw process, typically 10%–30%, although complete loss or damage is possible (45–47) (Table 1).

### Ejaculatory Duct Obstruction (EDO)

EDO is rare. When the ducts are obstructed in the urethra, transurethral resection of the ejaculatory ducts (TURED) relieves the obstruction where the distal ejaculatory duct terminates in the prostatic urethra (48). Resection may be guided by synchronous transurethral ultrasound (TUUS) to confirm obstruction location and avoid rectal injury (49). Relief of obstruction can often be seen immediately when cloudy fluid drains immediately on resection. Alternatively, seminal vesicle methylene blue chromotubation will release blue fluid upon resection (50). Sperm returns to the ejaculate in up to 70% (30/46) of men, and 20% (9/46) of couples will achieve unassisted pregnancy thereafter. Complications occur in up to 20% of patients and include urinary tract infection, epididymitis, hematuria, hematospermia, or watery ejaculate due to reflux of urine into the now widely patent ducts (51,52). When a midline cyst, not in connection with the ejaculatory duct system, causes the obstruction, aspiration of the cyst can relieve the blockage. This relief may be temporary or permanent and can be verified on a semen analysis soon

after the aspiration. Any sperm in the aspirated fluid can be cryopreserved if needed and suggests the need for resection.

Alternative therapies for EDO include endoscopic laser-assisted resection of the ducts, seminal vesicostomy or balloon dilation, or even antegrade seminal vesicle lavage to remove debris or calculi (53–55).

### TREATMENT SELECTION

#### Vasectomy Reversal vs. IVF/ICSI: Therapeutic Options

Both microsurgical reconstruction and sperm retrieval combined with IVF/ICSI can be effective treatments for infertility due to obstructive azoospermia. Several factors affect the decision; ultimately the choice must be based on the needs and preferences of the individual couple. Success rates and costs often are the factors weighted most heavily, but other considerations are relevant including risks of complications and the baseline fertility of each partner.

In experienced hands, microsurgical vasectomy reversal results in return of sperm to the ejaculate in 46.7% (n=30) to 98.1% (n=50) of patients (mean of 89%, N=6,633) (16), and up to 65%–75% of couples may achieve pregnancy without ART (12). Between 40% (n=14/35) (27) and 46% (n=6/13) (20) of couples may conceive naturally after bilateral end-to-side vasoepididymostomy.

The technique of sperm retrieval and the source of sperm (testis, epididymis, or vas deferens) have no significant effect on pregnancy rates achieved with IVF/ICSI (56). All methods generally provide sufficient numbers of viable sperm for ICSI and often also for cryopreservation. As long as viable sperm can be retrieved, neither the duration of obstruction (57, 58) nor the motility of the sperm affect the outcomes achieved with IVF/ICSI (59). When surgically retrieved frozen epididymal or testicular sperm are used for ICSI, fertilization rates range between 45% (n=1,335) and 74% (n=1,255) per

injected oocyte (35, 56, 60–62). Fertilization rates for fresh epididymal or testicular sperm are similar to frozen and range from 48% (n=242) to 72% (n=1,429) (35, 36, 56, 60), as with fertilization rates from two small studies of testicular sperm which did not specify fresh or frozen, 59% (N=21) to 81% (N=118) (56, 63). Clinical pregnancy rates from epididymal and testicular sperm (fresh and frozen) range from 22% (n=9 cycles) to 74% (n=24 cycles), with the largest reported groups from frozen epididymal and frozen epididymal sources, 24% (N=100 cycles) and 40% (n=145 cycles), respectively (36, 45, 64, 65). Delivery rates are less reported and range from 19.5% (testicular, n=41) to as high as 70.4% (epididymal, n=27) (64, 66).

### Impact of Female Factors

Clinical pregnancy and live-birth rates following vasectomy reversal or with sperm retrieval and IVF/ICSI are significantly impacted by characteristics of the female partner that correlate with female fertility. Outcomes are better in couples with the same female partner as before vasectomy. This advantage persists across the spectrum of female age and despite the duration of the obstructive interval (67).

Female age has been an independent predictor of success for both vasectomy reversal, with a younger age associated with better outcomes (68, 69). Studies find post-reversal pregnancy rates significantly worse when performed in men whose partners were 40 years or older (~15%) (70, 71), compared with under 39 years of age (55.7%) (71). Even after long obstructive intervals, female age is still the most profound predictor of success. In men with obstructive intervals greater than 15 years, the live-birth rates decreased with spousal age over 35. For spousal age <30 years, 30–35 years, 36–40 years, and >40 years, live-birth rates were 45% (n=22), 49% (n=39), 29% (n=29), and 14% (n=14), respectively (72).

An analysis of the Society for Assisted Reproductive Technology (SART) data demonstrates that among women undergoing ICSI/IVF for a male factor (a category that would include vasectomy), live-birth rates drop steadily with maternal age. The live-birth rate in 2014 varied by age for women under age 35 (48.7%), 35–37 (38.4%), 38–40 (24.3%), 41–42 (12.3%), and >42 years (3.8%), respectively (73). Ovarian reserve testing can help to select which option to choose when the female partner is age 34–40 when ART may shorten the time to pregnancy compared with vasectomy reversal, or in the 40 years-and-above population when ART success is diminished (74). The average time to pregnancy after a successful microsurgical vasectomy reversal is 12 months, although this may be impacted by female age (12). Consequently, sperm retrieval and IVF/ICSI may be the better option when the female partner is older and the window of opportunity to conceive is short. However, even with ART, fewer than 20% of women may be expected to achieve a successful pregnancy after age 40 (73). Sperm retrieval and IVF/ICSI also may be preferable to vasectomy reversal for couples having coexisting female infertility factors. When the female partner has significant tubal disease or a previous tubal sterilization procedure and both partners would require recon-

structive microsurgery, sperm retrieval with IVF/ICSI is the obvious choice.

### Secondary Male Infertility

A small proportion of men presenting for vasectomy reversal may have secondary infertility due to factors such as long-standing varicoceles, or internal injury, trauma, surgery, radiation, chemotherapy, or other spermatotoxic medication regimens since their last successful conception. For these men, the restorative potential of microsurgical reconstruction is compromised but sperm retrieval may still be possible, making IVF/ICSI the appropriate choice. Some men will be exposed to exogenous testosterone, which typically suppresses sperm production and creates a condition of nonobstructive azoospermia that is typically reversible but may take months to recover once testosterone is withheld. Gonadotropin support has been demonstrated to reverse this suppression successfully and allow for similar success rates of vasectomy reversal as unaffected men (75).

### Cost-effectiveness Comparisons

Microsurgical vasovasostomy (76) and vasoepididymostomy (77, 78) may be more cost-effective than sperm retrieval and IVF/ICSI, particularly for couples who hope to conceive more than a single pregnancy, because a successful reconstruction allows the couple to conceive naturally without further intervention. Even a repeated attempt at microsurgical reconstruction may be less costly than sperm retrieval and IVF/ICSI (79). Over the past several years, studies of cost-effectiveness have consistently demonstrated lower costs with microsurgical reconstruction. Such analyses include expected patency rates of at least 79% following reconstructive surgery, obstructive intervals, maternal age, and increased indirect costs from factors such as multiple conception seen with IVF/ICSI (76, 77, 80–84).

Advanced maternal age (above 40 years) combined with diminishing ovarian reserve typically moves the cost-effectiveness advantage toward IVF/ICSI, but as ovarian reserve drops below a threshold of advantage with ART, vasectomy reversal may again be more cost-effective due to the ability to attempt conception repeatedly without assistance (85). As the proficiency with ART has increased, live-birth outcomes with IVF/ICSI have improved and multiple gestations have decreased making ART more cost-effective. However, this cost advantage may be offset by expenses associated with the increased use of preimplantation genetic testing for aneuploidy (PGT-A). In circumstances of parental genetic-carrier states where the birth outcome may be seriously impacted, the indirect costs push the decision easily toward IVF/ICSI.

Ultimately, the results achieved with microsurgical reconstruction must be compared directly with those achieved with ART in the individual center. Furthermore, certain factors including the man's desire to maintain his vasal contraception or the couple's desire for having (an)other child(ren) may override financial calculations. Ideally, the final choice of treatment will be made by a well-informed couple

in consultation with the reproductive specialists caring for both partners.

For some individuals, the costs associated with retrieval of sperm and ART may be a barrier to treatment. In those cases, adoption and the use of donor sperm are other options that should be presented and discussed.

## SUMMARY

- Infertility due to obstructive azoospermia may be treated effectively by surgical reconstruction or by retrieval of sperm from the epididymis or testis, followed by IVF/ICSI.
- Sperm-retrieval techniques in general are minimally invasive and have a low complication risk. Successful sperm retrieval should be the expected outcome in the vast majority of obstructed men.

## CONCLUSIONS

- When obstructive azoospermia results from a vasectomy and there are no coexisting female infertility factors, microsurgical reconstruction of the reproductive tract is generally preferred over sperm retrieval and IVF/ICSI.
- Sperm retrieval with IVF/ICSI is generally the best choice of treatment for obstructive azoospermia when [1] the female partner has rapidly diminishing ovarian reserve due to age or other conditions, [2] there are coexisting female infertility factors that require IVF, [3] there is a concomitant secondary male infertility condition in the male partner, and [4] the likelihood for success with sperm retrieval/ICSI is greater than with surgical treatment.

**Acknowledgments:** This report was developed under the direction of the Practice Committee of the American Society for Reproductive Medicine as a service to its members and other practicing clinicians. Although this document reflects appropriate management of a problem encountered in the practice of reproductive medicine, it is not intended to be the only approved standard of practice or to dictate an exclusive course of treatment. Other plans of management may be appropriate, taking into account the needs of the individual patient, available resources, and institutional or clinical practice limitations. The Practice Committee and the Board of Directors of the American Society for Reproductive Medicine have approved this report.

This document was reviewed by ASRM members, and their input was considered in the preparation of the final document. The Practice Committee acknowledges the special contribution of Jay Sandlow, M.D.; Christopher M. Deibert, M.D.; Akanksha Mehta, M.D.; and Aaron Spitz, M.D. in the preparation of this document. The following members of the ASRM Practice Committee participated in the development of this document. All Committee members disclosed commercial and financial relationships with manufacturers or distributors of goods or services used to treat patients. Members of the Committee who were found to have conflicts of interest based on the relationships disclosed did not participate in the discussion or development of this document.

Alan Penzias, M.D.; Kristin Bendikson, M.D.; Samantha

Butts, M.D., M.S.C.E.; Christos Coutifaris, M.D., Ph.D.; Tommaso Falcone, M.D.; Susan Gitlin, Ph.D.; Clarisa Gracia, M.D., M.S.C.E.; Karl Hansen, M.D., Ph.D.; Sangita Jindal, Ph.D.; Suleena Kalra, M.D., M.S.C.E.; Jennifer Mersereau, M.D.; Randall Odem, M.D.; Robert Rebar, M.D.; Richard Rein-dollar, M.D.; Mitchell Rosen, M.D.; Jay Sandlow, M.D.; Peter Schlegel, M.D.; Dale Stovall, M.D.; Michael Vernon, Ph.D.

## REFERENCES

1. Jarow JP, Espeland MA, Lipshultz LI. Evaluation of the azoospermic patient. *J Urol* 1989;142:62–5.
2. Practice Committee of the American Society for Reproductive Medicine. The management of nonobstructive azoospermia: a committee opinion. *Fertil Steril* 2018;109:777–82.
3. Silber SJ. Perfect anatomical reconstruction of vas deferens with a new microscopic surgical technique. *Fertil Steril* 1977;28:72–7.
4. Owen ER. Microsurgical vasovasostomy: a reliable vasectomy reversal. *Aust N Z J Surg* 1977;47:305–9.
5. Kim HH, Goldstein M. History of vasectomy reversal. *Urol Clin North Am* 2009;36:359–73.
6. Nangia AK, Myles JL, Thomas AJ. Vasectomy reversal for the post-vasectomy pain syndrome: a clinical and histological evaluation. *J Urol* 2000;164:1939–42.
7. Belker AM. Infrapubic incision for specific vasectomy reversal situations. *Urology* 1988;32:413–5.
8. Silber SJ. Pregnancy after vasovasostomy for vasectomy reversal: a study of factors affecting long-term return of fertility in 282 patients followed for 10 years. *Hum Reprod* 1989;4:318–22.
9. Witt MA, Heron S, Lipshultz LI. The post-vasectomy length of the testicular vasa remnant: a predictor of surgical outcome in microscopic vasectomy reversal. *J Urol* 1994;151:892–4.
10. Bolduc S, Fischer MA, Deceuninck G, Thabet M. Factors predicting overall success: a review of 747 microsurgical vasovasostomies. *Can Urol Assoc J* 2007;1:388–94.
11. Patel SR, Sigman M. Comparison of outcomes of vasovasostomy performed in the convoluted and straight vas deferens. *J Urol* 2008;179:256–9.
12. Belker AM, Thomas AJ Jr, Fuchs EF, Konnak JW, Sharlip ID. Results of 1,469 microsurgical vasectomy reversals by the Vasovasostomy Study Group. *J Urol* 1991;145:505–11.
13. Sigman M. The relationship between intravasal sperm quality and patency rates after vasovasostomy. *J Urol* 2004;171:307–9.
14. Hsieh ML, Huang HC, Chen Y, Huang ST, Chang PL. Loupe-assisted vs microsurgical technique for modified one-layer vasovasostomy: is the micro-surgery really better? *BJU Int* 2005;96:864–6.
15. Safarinejad MR, Lashkari MH, Asgari SA, Farshi A, Babaei AR. Comparison of macroscopic one-layer over number 1 nylon suture vasovasostomy with the standard two-layer microsurgical procedure. *Hum Fertil (Camb)* 2013;16:194–9.
16. Herrel LA, Goodman M, Goldstein M, Hsiao W. Outcomes of microsurgical vasovasostomy for vasectomy reversal: a meta-analysis and systematic review. *Urology* 2015;85:819–25.
17. Thomas AJ Jr. Vasoepididymostomy. *Urol Clin North Am* 1987;14:527–38.
18. Berger RE. Triangulation end-to-side vasoepididymostomy. *J Urol* 1998;159:1951–3.
19. Marmar JL. Modified vasoepididymostomy with simultaneous double needle placement, tubulotomy and tubular invagination. *J Urol* 2000;163:483–6.
20. Schiff J, Chan P, Li PS, Finkelberg S, Goldstein M. Outcome and late failures compared in 4 techniques of microsurgical vasoepididymostomy in 153 consecutive men. *J Urol* 2005;174:651–5, quiz 801.
21. Parekattil SJ, Gudeloglu A, Brahmabhatt J, Wharton J, Priola KB. Robotic assisted versus pure microsurgical vasectomy reversal: technique and prospective database control trial. *J Reconstr Microsurg* 2012;28:435–44.
22. Barazani Y, Kaouk J, Sabanegh ES Jr. Robotic intra-abdominal vasectomy reversal: a new approach to a difficult problem. *Can Urol Assoc J* 2014;8:E439–41.

23. Trost L, Parekattil S, Wang J, Hellstrom WJ. Intracorporeal robot-assisted microsurgical vasovasostomy for the treatment of bilateral vasal obstruction occurring following bilateral inguinal hernia repairs with mesh placement. *J Urol* 2014;191:1120–5.
24. Matthews GJ, Schlegel PN, Goldstein M. Patency following microsurgical vasoepididymostomy and vasovasostomy: temporal considerations. *J Urol* 1995;154:2070–3.
25. Jarow JP, Sigman M, Buch JP, Oates RD. Delayed appearance of sperm after end-to-side vasoepididymostomy. *J Urol* 1995;153:1156–8.
26. Boorjian S, Lipkin M, Goldstein M. The impact of obstructive interval and sperm granuloma on outcome of vasectomy reversal. *J Urol* 2004;171:304–6.
27. Chan PT, Brandell RA, Goldstein M. Prospective analysis of outcomes after microsurgical intussusception vasoepididymostomy. *BJU Int* 2005;96:598–601.
28. Hollingsworth MR, Sandlow JJ, Schrepferman CG, Brannigan RE, Kolettis PN. Repeat vasectomy reversal yields high success rates. *Fertil Steril* 2007;88:217–9.
29. Hernandez J, Sabanegh ES. Repeat vasectomy reversal after initial failure: overall results and predictors for success. *J Urol* 1999;161:1153–6.
30. Paick JS, Park JY, Park DW, Park K, Son H, Kim SW. Microsurgical vasovasostomy after failed vasovasostomy. *J Urol* 2003;169:1052–5.
31. Pasqualotto FF, Agarwal A, Srivastava M, Nelson DR, Thomas AJ Jr. Fertility outcome after repeat vasoepididymostomy. *J Urol* 1999;162:1626–8.
32. Silber SJ, Nagy ZP, Liu J, Godoy H, Devroey P, Van Steirteghem AC. Conventional in-vitro fertilization versus intracytoplasmic sperm injection for patients requiring microsurgical sperm aspiration. *Hum Reprod* 1994;9:1705–9.
33. Schlegel PN, Palermo GD, Alikani M, Adler A, Reing AM, Cohen J, et al. Micropuncture retrieval of epididymal sperm with in vitro fertilization: importance of in vitro micromanipulation techniques. *Urology* 1995;46:238–41.
34. Cayan S, Lee D, Conaghan J, Givens CA, Ryan IP, Schriock ED, et al. A comparison of ICSI outcomes with fresh and cryopreserved epididymal spermatozoa from the same couples. *Hum Reprod* 2001;16:495–9.
35. Nagy Z, Liu J, Cecile J, Silber S, Devroey P, Van Steirteghem A. Using ejaculated, fresh, and frozen-thawed epididymal and testicular spermatozoa gives rise to comparable results after intracytoplasmic sperm injection. *Fertil Steril* 1995;63:808–15.
36. Tournaye H, Merdad T, Silber S, Joris H, Verheyen G, Devroey P, et al. No differences in outcome after intracytoplasmic sperm injection with fresh or with frozen-thawed epididymal spermatozoa. *Hum Reprod* 1999;14:90–5.
37. Oldereid NB, Haneyik HI, Bakkevig I, Romundstad LB, Magnus Ø, Hazekamp J, et al. Pregnancy outcome according to male diagnosis after ICSI with non-ejaculated sperm compared with ejaculated sperm controls. *Reprod Biomed Online* 2014;29:417–23.
38. Schlegel PN, Berkeley AS, Goldstein M, Cohen J, Alikani M, Adler A, et al. Epididymal micropuncture with in vitro fertilization and oocyte micromanipulation for the treatment of unreconstructable obstructive azoospermia. *Fertil Steril* 1994;61:895–901.
39. Tournaye H, Devroey P, Liu J, Nagy Z, Lissens W, Van Steirteghem A. Microsurgical epididymal sperm aspiration and intracytoplasmic sperm injection: a new effective approach to infertility as a result of congenital bilateral absence of the vas deferens. *Fertil Steril* 1994;61:1045–51.
40. Practice Committee of American Society for Reproductive Medicine. Sperm retrieval for obstructive azoospermia. *Fertil Steril* 2008;90:S213–8.
41. Ubaldi F, Camus M, Tournaye H, Clasen K, Nagy Z, Smitz J, et al. Results of microsurgical epididymal sperm aspiration (MESA) and testicular sperm extraction (TESE) in azoospermic men using intracytoplasmic sperm injection (ICSI). *Andrologia* 1996;28:71–5.
42. van Wely M, Barbey N, Meissner A, Repping S, Silber SJ. Live birth rates after MESA or TESE in men with obstructive azoospermia: is there a difference? *Hum Reprod* 2015;30:761–6.
43. Wood S, Thomas K, Sephton V, Troup S, Kingsland C, Lewis-Jones I. Postoperative pain, complications, and satisfaction rates in patients who undergo surgical sperm retrieval. *Fertil Steril* 2003;79:56–62.
44. Jensen CF, Ohl DA, Hiner MR, Fode M, Shah T, Smith GD, et al. Multiple needle-pass percutaneous testicular sperm aspiration as first-line treatment in azoospermic men. *Andrology* 2016;4:257–62.
45. Garg T, LaRosa C, Strawn E, Robb P, Sandlow JJ. Outcomes after testicular aspiration and testicular tissue cryopreservation for obstructive azoospermia and ejaculatory dysfunction. *J Urol* 2008;180:2577–80.
46. Schiewe MC, Rothman C, Spitz A, Werthman PE, Zeitlin SI, Anderson RE. Validation-verification of a highly effective, practical human testicular tissue in vitro culture-cryopreservation procedure aimed to optimize pre-freeze and post-thaw motility. *J Assist Reprod Genet* 2016;33:519–28.
47. Ohlander S, Hotaling J, Kirshenbaum E, Niederberger C, Eisenberg ML. Impact of fresh versus cryopreserved testicular sperm upon intracytoplasmic sperm injection pregnancy outcomes in men with azoospermia due to spermatogenic dysfunction: a meta-analysis. *Fertil Steril* 2014;101:344–9.
48. Farley S, Barnes R. Stenosis of ejaculatory ducts treated by endoscopic resection. *J Urol* 1973;109:664–6.
49. Modgil V, Rai S, Ralph DJ, Muneer A. An update on the diagnosis and management of ejaculatory duct obstruction. *Nat Rev Urol* 2016;13:13–20.
50. Purohit RS, Wu DS, Shinohara K, Turek PJ. A prospective comparison of 3 diagnostic methods to evaluate ejaculatory duct obstruction. *J Urol* 2004;171:232–5, discussion 235–6.
51. Turek PJ, Magana JO, Lipshultz LI. Semen parameters before and after transurethral surgery for ejaculatory duct obstruction. *J Urol* 1996;155:1291–3.
52. Smith JF, Walsh TJ, Turek PJ. Ejaculatory duct obstruction. *Urol Clin North Am* 2008;35:221–7.
53. Halpern EJ, Hirsch IH. Sonographically guided transurethral laser incision of a Müllerian duct cyst for treatment of ejaculatory duct obstruction. *AJR Am J Roentgenol* 2000;175:777–8.
54. Colpi GM, Negri L, Patrizio P, Pardi G. Fertility restoration by seminal tract washout in ejaculatory duct obstruction. *J Urol* 1995;153:1948–50.
55. Xu B, Niu X, Wang Z, Li P, Qin C, Li J, et al. Novel methods for the diagnosis and treatment of ejaculatory duct obstruction. *BJU Int* 2011;108:263–6.
56. Palermo GD, Schlegel PN, Hariprashad JJ, Ergün B, Mielnik A, Zaninovic N, et al. Fertilization and pregnancy outcome with intracytoplasmic sperm injection for azoospermic men. *Hum Reprod* 1999;14:741–8.
57. Sukcharoen N, Sithipravej T, Promviengchai S, Chinpilas V, Boonkasemsanti W. No differences in outcome of surgical sperm retrieval with intracytoplasmic sperm injection at different intervals after vasectomy. *Fertil Steril* 2000;74:174–5.
58. Nicopoulos JD, Gilling-Smith C, Almeida PA, Ramsay JW. Effect of time since vasectomy and maternal age on intracytoplasmic sperm injection success in men with obstructive azoospermia after vasectomy. *Fertil Steril* 2004;82:367–73.
59. Moghadam KK, Nett R, Robins JC, Thomas MA, Awadalla SG, Scheiber MD, et al. The motility of epididymal or testicular spermatozoa does not directly affect IVF/ICSI pregnancy outcomes. *J Androl* 2005;26:619–23.
60. Habermann H, Seo R, Cieslak J, Niederberger C, Prins GS, Ross L. In vitro fertilization outcomes after intracytoplasmic sperm injection with fresh or frozen-thawed testicular spermatozoa. *Fertil Steril* 2000;73:955–60.
61. Mercan R, Urman B, Alatas C, Aksoy S, Nuhoglu A, Isiklar A, et al. Outcome of testicular sperm retrieval procedures in non-obstructive azoospermia: percutaneous aspiration versus open biopsy. *Hum Reprod* 2000;15:1548–51.
62. Küpker W, Schlegel PN, Al-Hasani S, Fornara P, Johannisson R, Sandmann J, et al. Use of frozen-thawed testicular sperm for intracytoplasmic sperm injection. *Fertil Steril* 2000;73:453–8.
63. Gil-Salom M, Meinguez Y, Rubio C, De los Santos MJ, Remohí J, Pellicer A. Efficacy of intracytoplasmic sperm injection using testicular spermatozoa. *Hum Reprod* 1995;10:3166–70.
64. Anger JT, Wang GJ, Boorjian SA, Goldstein M. Sperm cryopreservation and in vitro fertilization/intracytoplasmic sperm injection in men with congenital bilateral absence of the vas deferens: a success story. *Fertil Steril* 2004;82:1452–4.
65. Sharma RK, Padron OF, Thomas AJ Jr, Agarwal A. Factors associated with the quality before freezing and after thawing of sperm obtained by microsurgical epididymal aspiration. *Fertil Steril* 1997;68:626–31.
66. Heidenreich A, Altmann P, Engelmann UH. Microsurgical vasovasostomy versus microsurgical epididymal sperm aspiration/testicular extraction of sperm combined with intracytoplasmic sperm injection. A cost-benefit analysis. *Eur Urol* 2000;37:609–14.

67. Ostrowski KA, Polackwich S, Kent J, Conlin MJ, Hedges JC, Fuchs EF. Higher outcomes of vasectomy reversal in men with the same female partner as before vasectomy. *J Urol* 2015;193:245–7.
68. Reddy UM, Wapner RJ, Rebar RW, Tasca RJ. Infertility, assisted reproductive technology, and adverse pregnancy outcomes: executive summary of a National Institute of Child Health and Human Development workshop. *Obstet Gynecol* 2007;109:967–77.
69. Hull MG, Fleming CF, Hughes AO, McDermott A. The age-related decline in female fecundity: a quantitative controlled study of implanting capacity and survival of individual embryos after in vitro fertilization. *Fertil Steril* 1996;65:783–90.
70. Hinz S, Rais-Bahrami S, Kempkensteffen C, Weiske WH, Schrader M, Magheli A. Fertility rates following vasectomy reversal: importance of age of the female partner. *Urol Int* 2008;81:416–20.
71. Gerrard ER Jr, Sandlow JI, Oster RA, Burns JR, Box LC, Kolettis PN. Effect of female partner age on pregnancy rates after vasectomy reversal. *Fertil Steril* 2007;87:1340–4.
72. Fuchs EF, Burt RA. Vasectomy reversal performed 15 years or more after vasectomy: correlation of pregnancy outcome with partner age and with pregnancy results of in vitro fertilization with intracytoplasmic sperm injection. *Fertil Steril* 2002;77:516–9.
73. Society for Assisted Reproductive Technology. SART CORS online. Clinic summary report: all SART member clinics, IVF success rates. Available at: [https://www.sartcorsonline.com/rptCSR\\_PublicMultYear](https://www.sartcorsonline.com/rptCSR_PublicMultYear). Accessed January 24, 2019.
74. Shridharani A, Sandlow JI. Vasectomy reversal versus IVF with sperm retrieval: which is better? *Curr Opin Urol* 2010;20:503–9.
75. Coward RM, Mata DA, Smith RP, Kovac JR, Lipshultz LI. Vasectomy reversal outcomes in men previously on testosterone supplementation therapy. *Urology* 2014;84:1335–41.
76. Pavlovich CP, Schlegel PN. Fertility options after vasectomy: a cost-effectiveness analysis. *Fertil Steril* 1997;67:133–41.
77. Kolettis PN, Thomas AJ Jr. Vasoepididymostomy for vasectomy reversal: a critical assessment in the era of intracytoplasmic sperm injection. *J Urol* 1997;158:467–70.
78. Paick JS, Hong SK, Yun JM, Kim SW. Microsurgical single tubular epididymo-vasostomy: assessment in the era of intracytoplasmic sperm injection. *Fertil Steril* 2000;74:920–4.
79. Donovan JF Jr, DiBaise M, Sparks AE, Kessler J, Sandlow JI. Comparison of microscopic epididymal sperm aspiration and intracytoplasmic sperm injection/in vitro fertilization with repeat microscopic reconstruction following vasectomy: is second attempt vas reversal worth the effort? *Hum Reprod* 1998;13:387–93.
80. Lee R, Li PS, Goldstein M, Tanrikut C, Schattman G, Schlegel PN. A decision analysis of treatments for obstructive azoospermia. *Hum Reprod* 2008;23:2043–9.
81. Robb P, Sandlow JI. Cost-effectiveness of vasectomy reversal. *Urol Clin North Am* 2009;36:391–6.
82. Meng MV, Greene KL, Turek PJ. Surgery or assisted reproduction? A decision analysis of treatment costs in male infertility. *J Urol* 2005;174:1926–31.
83. Hsieh MH, Meng MV, Turek PJ. Markov modeling of vasectomy reversal and ART for infertility: how do obstructive interval and female partner age influence cost effectiveness? *Fertil Steril* 2007;88:840–6.
84. Lee R, Li PS, Schlegel PN, Goldstein M. Reassessing reconstruction in the management of obstructive azoospermia: reconstruction or sperm acquisition? *Urol Clin North Am* 2008;35:289–301.
85. Sandlow J. Vasectomy reversal versus IVF with sperm retrieval; which better? *Curr Opin Urology* 2010;20:503–9.