Vasectomy reversal

MRA Young, CJH Logan

Accepted 8 August 1989.

SUMMARY

A small personal series of vasovasostomies and a review of the literature on this subject are presented. The importance is stressed of good counselling before vasectomy is undertaken in the first place.

INTRODUCTION

The popularity of vasectomy is increasing, but so too is the rate of divorce and remarriage. There is also more awareness of the possibility that this operation can be reversed.

METHODS

Thirteen vasovasostomies were performed by one of the authors (HL) between 1978–1988. Follow up information was gained by personal interview, seminal analysis, post-reversal haematological tests and case records. Three patients were not contactable due to change in address.

All operations involved bilateral identification and limited mobilization of the previously divided vas deferens. A 3/0 gauge nylon double ended straight needle stent suture was advanced up the lumen of the proximal and distal vas for 1.5 cms and then brought through their walls. Four 6/0 nylon seromuscular sutures completed the anastomosis. The stent sutures were brought through the skin and ligated in loose loops. Patients had bed rest for two days and stents were removed on the tenth day. Seminal analyses were performed postoperatively at 1-2 months and again between 3-5 months.

In addition to seminal analysis at recent review, the Tray agglutination test ¹ assessed sperm and seminal antibodies, and a complement preparation was used for sperm immobilising antibodies (positive control from Wellcome Laboratories Ltd).

RESULTS

Clinical, operative and laboratory data on the 13 patients are shown in the Table. The average age at vasectomy was $27 \cdot 6$ years (range $21 \cdot 5$ to 38 years) and at reversal $33 \cdot 3$ years (range 23 - 49 years). Four patients had vasectomy at 24 years of age or under. Remarriage following divorce was the basis for reversal in ten patients, two requested more children (vasectomy at 21 years and 24 years), and one widower was remarrying.

The Ulster Hospital, Dundonald, Belfast BT16 0RH.

MRA Young, FRCS, Registrar.

C J H Logan, FRCS, Consultant Surgeon.

Correspondence to Mr Logan.

TABLE
Clinical, operative and laboratory data on 13 patients

No.	1	2	3	4	5	6	7	8	9	10	11	12	13
Age at vasectomy	24	26	38	23	21	28	33	32	26	30	27	22	28
Age at reversal	27	30	49	30	23	31	43	36	30	36	29	36	33
Interval (years)	3	4	11	7	1	3	10	4	4	6	2	14	5
Operative findings													
Poor lumen	_	+	_	_	_	_	_	+	_	_	_	_	_
Convoluted	_	. +	_	+	-	-	-	+	-	-	-	-	_
Laboratory data Sperm counts (X10 ⁶ /ml)	i												
Initial	7	nil	20	4.8	0.3	84		nil	24			138	nil
Recent	10	nil	nil	10	8	120	nil	nil		•	9		nil
Tray agglutination test	– ve	– ve	– ve	– ve	– ve	– ve	1:128	– ve	•	•	– ve		– ve
Successful pregnancy post				_				_			_		
operative	0	0	N/A	0	1	0	0	0	•	•	3	•	N/A

(N/A not applicable)

Three patients, one of whom had no post-operative data, could not be contacted for the recent analysis and pregnancy information. Twelve patients had a sperm count performed initially and/or on review. Of these, eight had a positive count giving a patency rate of 66%.

Eight patients had complete data; of these three were azospermic throughout, and of the remaining five patients four had an increase in their sperm count with time. The remaining patient developed prostatitis in the interim between the tests. Two of the four azospermic patients had a poor lumen at vasovasostomy and had had vasectomy in the convoluted portion of the vas deferens. One had post-operative granuloma formation and the other had macroscopic normal anatomy.

It is difficult to assess the ultimate pregnancy rate as three patients were not contacted and two are hoping to conceive in the future. The wives of two patients have become pregnant, one before he brought any specimens for examination. Their sperm counts were eight and nine million/ml. Only one serum Tray agglutination test was positive (1:128). This patient had no sperm production, a good lumen but developed a post-operative granuloma. No immobilizing or seminal antibodies were detected.

DISCUSSION

The pathophysiological events following vasectomy are not entirely understood but back-pressure induced congestion and dilatation within the testes and

[©] The Ulster Medical Society, 1989.

proximal vas deferens results in decreased patency and subsequent sperm counts.^{2, 3} This problem is more significant as the interval between vasectomy and its reversal increases. Obstruction of more than five years is regarded as unfavourable,⁴ but the patient's age is not considered relevant for success.

Despite a successful anatomical vasovasostomy, back pressure may cause epididymal sperm granuloma following tubal rupture with resultant functional failure. However, sperm granuloma, created by a continuous leakage at the vasectomy site have a beneficial pressure-releasing effect, with improved sperm quality and less dilation.^{2, 5} Sperm granuloma should be proven histologically as they may be confused with suture granuloma.

Controversy exists as to the optimum method of anastomosis. Microscopic, macroscopic, and stented anastomoses are techniques advocated for vasovasostomy or vaso-epididymostomy. The general principle in all techniques includes limited mobilization to minimise disturbance of the blood supply, longitudinal incisions to preserve the cremasteric peristalsis and an accurate leakproof anastomosis to reduce subsequent scarring. Favourable results of 83% sperm present and 55% pregnancy rates are reported for externalized nylon stents. Similar results are documented for +2.5 or +3.5 loupe magnification of stenting, while Denton reports 65% pregnancy rates using a stented macroscopic anastomosis and a 61% rate without stents. Canine experiments have shown that vas obstruction occurs at the stent exit and not at the anastomosis.

Macroscopic techniques without stents have varying and less favourable results. This ranges from a 19% pregnancy rate in a questionnaire of urologists (1630 patients), 10 a 39% pregnancy rate 11 for a two layer anastomosis, to a 57% pregnancy rate 12 using a +2.5 loupe magnification of a one layer full thickness anastomosis. Amelar and Dubin 13 were disappointed with a 33% pregnancy rate and changed to a micro-surgical technique with 53% success rate.

An improved pregnancy rate was observed with the advent of the microscopic technique, advocated by Sibler ¹⁴ in 1975, with a pregnancy rate of 76%. Owens, ¹⁵ had comparable results (72%) from this two layer microscopic technique. Belker ¹⁴ reported a 62% pregnancy rate using a × 10 to × 20 operating microscope (10/0 mucosal and 9/0 muscular anastomosis). Generally one and two layer microscopic anastomoses have similar results: Sharlip ¹⁶ had 50% pregnancies with two layers and 52% with one layer anastomoses while Lee ³ had 52% and 50% for the same techniques. While microscopic anastomoses attain optimal anatomical results, considerable practice is required to master the operating microscope. Some feel this expertise does not necessarily impart improved results in view of reported pregnancy rates of greater than 50% for any technique and that results are more dependent on the surgeon than on technique. ¹⁷

Absence of sperm heads or a creamy paste consistency to the seminal fluid at intra-operative examination is an unfavourable prognostic feature. Better grades of fluid are observed when sperm granuloma are present. Half of the patients azospermic at operation have a reasonable recovery. Generally, sperm counts and motility improve with time though re-exploration can be successful if secondary scarring becomes evident by diminishing sperm counts. Vasectomies performed in the convoluted portion of the vas deferens or when a long segment has been excised can make the anastomosis technically difficult.

Technique may explain patency, but a discrepancy exists between patency and pregnancy rates. Sperm reappearance is therefore not necessarily an index of fertility. An immunological basis may explain this. Sperm antibodies occur in two thirds of vasectomized men, especially when sperm granuloma are present. Most studies correlate high antibody titres with infertility. The Tray agalutinating test detects head to head sperm agglutination (IgM and IgG) which is common in vasectomy patients and rare in naturally infertile men (prostatic IqA).¹⁸ Despite diminished fertility with serum agalutinating antibody titres of 1:64, titres of 1:256 correlate better with infertility, 18, 19 but 25% of patients still conceive with these high titres. Semen agglutinating activity and immobilizing antibodies are only detected when there are high titres of serum agglutinating antibody. Seminal agalutinating activity significantly increases after vasectomy reversal 18, 19, 20 whereas serum levels remain relatively unaltered. Pregnancy rates are reduced with seminal titres of > 1:8 (65% to 7% 20 and 85% to 14% 19 pregnancy rates for negative and positive seminal agglutinating activity respectively). Hence seminal agglutinating activity is closely correlated with infertility. Immunosuppression by steroids is claimed to have beneficial effect in reducing antibody titres.7, 21

Most wives who become pregnant do so within two years. If not and the male partner appears fertile then the female's fertility should be questioned. In summary, mechanical failure results from a long interval, excessive vas excision, vasectomy in the convoluted portion, poor anastomosis, infection-induced fibrosis and anastomotic granuloma. Function failure with poor semen quality results from back pressure, injury to sympathetic fibres, sperm antibodies and low fertility of the wife.

This small study highlights the social background in our society and good prevasectomy counselling in Northern Ireland. Restricting vasectomy to those in their late twenties and over would further reduce reversal requests. Our study identifies no hindrance from a long period of obstruction. Only one patient had a positive antibody titre. Our technique appears correct as control samples were positive. Generally, post-reversal sperm counts were lower than would be regarded as normal but this appears to be no obstacle to conceiving. Operative findings of poor lumen, granuloma and anastomosis in the convoluted portion of the vas correlate best with poor results. It would seem reasonable practice to anastomose both vasa to double the chances of success. To reduce stent exit granuloma we suggest that a long length of stent is passed down the distal vas and brought out through one exit site on the proximal vas.

We are grateful to Mr George Greer for performing the serological analyses.

REFERENCES

- Jequier AM, Crich AJP. Seminal analysis. A practical guide. London: Blackwell Scientific Publications, 1986.
- 2. Sibler SJ. Sperm granuloma and reversibility of vasectomy. Lancet 1977; 2: 588-9.
- 3. Lee HY. A twenty year experience with vasovasostomy. J Urol 1986; 136: 413-5.
- 4. Belker AM. Vasectomy reversal. Urol Clin North Am 1987; 14: 155-66.
- 5. Belker AM, Konnak JW, Sharlip ID, Thomas AJ. Intraoperative observations during vasovasostomy in 334 patients. *J Urol* 1983; **129**: 524-7.
- © The Ulster Medical Society, 1989.

- Phadke GM, Phadke AG. Experiences in the re-anastomosis of the vas deferens. J Urol 1967; 97: 888-90.
- 7. Urquhart-Hay D. A low-power magnification technique for the re-anastomosis of the vas further results in a personal series of 125 patients. Aust NZ J Surg 1984; 54: 73-4.
- 8. Denton SE, Bohnert WW, Kurtz CW. Vasectomy reversal technique and results. *Arizona Med (Surg)* 1983; **40**: 33-6.
- Fernandes M, Shah KN, Draper JW. Vasovasostomy: improved microsurgical technique. J Urol 1968; 100: 763-6.
- Derrick FC, Yarbrough W, D'Agostino J. Vasovasostomy: results of questionnaire to members of the American Urological Association. J Urol 1973; 110: 556-7.
- 11. Middleton RG, Henderson D. Vas deferens re-anastomosis without splints and without magnification. *J Urol* 1978: **119**: 763-4.
- 12. Fallon B, Millar RK, Gerber WL. Non microscopic vasovasostomy. J Urol 1981; 126: 361-2.
- 13. Amelar RD, Dubin L. Vasectomy reversal. J Urol 1979; 121: 547-50.
- 14. Sibler SJ. Microsurgery in clinical urology. *Urol* 1975; **6**: 150-3.
- Owen ER. Microsurgical vasovasostomy: a reliable vasectomy, reversal. Aust NZ J Surg 1977;
 47: 305-9.
- Sharlip ID. Microsurgical vasovasostomy. A comparison of double and modified one layer anastomosis. American Urological Association Annual Meeting. 1985; Abst 696.
- 17. Fenster H, McLoughlin MG. Vasovasostomy is the microscope necessary? *Urol* 1981; 18: 60-4.
- Parslow JM, Royce MG, Kingscott MMB, Wallace DMA, Hendry WF. The effect of sperm antibodies on fertility after vasectomy reversal. Am J Reprod Immunol 1983; 3: 28-31.
- 19. Linnet L, Hjort T, Fogh-Andersen P. Association between failure to impregnate after vasovasostomy and sperm agglutinins in semen. *Lancet* 1981; 1: 117-9.
- Sutherland PD, Matson PC, Masters JRW, Pryor JP. Association between infertility following reversal of vasectomy and the presence of sperm agglutinating activity in semen. *Int J Androl* 1984; 7: 503-8.
- Mathur S, Baker ER, Williamson HO, Derrick FC, Teague KJ, Fudenberg HH. Clinical significance of sperm antibodies in infertility. Fertil Steril 1981; 36: 486-95.