

FEASIBILITY, COMPLICATIONS & SHORT TERM RESULTS OF VAGINAL APEX OR VAULT SUSPENSION WITH POSTERIOR IVS® (PRELIMINARY RESULTS ON 113 CASES).

Hypothesis/aims of the study

We performed a prospective clinical study to evaluate the feasibility, the complications and the results of the vaginal apex or vault suspension with the posterior IVS®.

Study design/materials and methods

From May 2002 to december 2003, 113 patients from 2 belgian centres have been treated for vaginal vault (v.v.) prolapse with posterior IVS®. All the patients complain about prolapse. The mean age is 63,2 years (33-83). The majority of the patients has previous pelvic operation: hysterectomy (n=59), cystocele (n=16) & rectocele (n=7) repair, no v.v. suspension.

Pre-operative examination shows 112 v.v. or uterine prolapse: 29 stage 1 (POPQ -8 à -1), 46 stage 2 (POPQ -1 à +1), 26 stage 3 (POPQ >+1) & 8 stage 4 (POPQ >+8), 1 stage 0 with elythro-rectocele stage 3 (POPQ >+1). The associated prolapses are: 20 cystocele stage 1 (POPQ -3 à -1), 28 stage 2 (POPQ -1 à +1), 44 stage 3 (POPQ >+1); 25 rectoceles stage 1 (POPQ -3 à -1), 47 stage 2 (POPQ -1 à +1), 29 stage 3 (POPQ >+1). Fifty four patients have GSI. Seventy three (65%) complain of urge incontinence &/or pollakiuria &/or nocturia; 45 (40%) has dyschesia.

The operative technique described by P. Papapetros modified by P. Von Théobald has been used in all cases: posterior colpotomy, dissection of the pararectal space to the ischial spine, finger-guided progression of the tunneler through the ischio-rectal space & the pelvic muscles, suture of the tape on the utero-sacral ligaments or the cervix. Spinal anesthesia is commonly performed (82%). In 109 cases, the posterior IVS® is associated to other vaginal surgery: 55 TOT/TVT, 6 cervix ablation, 27 hysterectomy, 78 cystocele & 76 rectocele repair with or without graft.

Results

No peroperative complication due to the IVS® occurred: no bladder or rectal perforation, no hemorrhage. During the hospital stay, we noticed 8 postoperative hematomas of the recto-vaginal or para-rectal space. Seven have been drained spontaneously, 1 under general anesthesia. One hundred & four patients (92%) have been clinically reviewed between 3 weeks & 20 months; 61% of them have a mean follow-up of 10 months. The post-operative results show a good suspension of the v.v. in 97% (n=101; POP-Q <-8). Two patients have early recurrence of v.v. prolapse (4 & 6 months) and one at 19 months. All the three have a POP-Q of <-1 to +1. No « de novo » dysuria, dyschesia, or overactive bladder symptoms occurred. One patient developed GSI. Five patients (5%) had wound complications: 1 granuloma and 4 erosions of the vaginal vault with partial rejection of the IVS® between 4 & 15 months. One patient developed a chronic cutaneo-vaginal fistula along the IVS® documented by fistulography. They all have been treated by ablation of the non-colonised tape and local desinfection. Four of the five erosions followed a post-operative hematoma but without predictive event during surgery. In all cases the v.v. suspension remained.

Conclusions

The treatment of the v.v. or uterine prolapse with posterior IVS® is a reproducible and efficient technique (97% of success) without peroperative complications. These results are comparable to the literature (). But postoperative complications as rejections of the tape (5%) and early recurrences (3%) are more frequent than in the others publications. These complications could be avoided. The type of the IVS® tape (polypropilene, multifilament) and the large dissection with risk of postoperative bleeding, are probably responsible of recurrences and tape infection and rejection. A monofilament or may be a biomaterial mesh, a smaller dissection with very careful hemostasis could probably improve the results.