

General (49)	1,5±1,8	3,6±2,2	1,5±4,1	8±1,2	-0,4±2	0±2,6	-3,2±2,4
Anterior (31)	1,4±1,7	3,6±1,7	0,4±3,6	7,9±1	-1,2±1,6	-0,9±2,1	-3,7±2,2
Apical (15)	2,2±1,3	4,7±1,8	6±1,8	8,3±1,5	0,9±2	1,7±2,9	-1,5±1,6
Posterior (3)	-2±1	-2,3±1,1	-4,3±1,5	7±1,7	2,6±1,1	3,6±0,5	-7±1,4

Surgical details:

Forty-three surgeries were under regional anaesthesia, the rest were under general anaesthesia. Perioperative data are shown in table 4.

Table 4

Perioperative Data	
Mean operating time±SD, range/mode (min)	50,2±19,3, 25-120/40
Mean Estimated blood loss±SD, range/mode (ml)	56,6±44,5, 10-200/50
Intraoperative complications	0
Concomitant vaginal surgeries	6 ¹
Median hospital stay±SD, range/mode (days)	2,6±2,99, 1-20/2

¹ One TOT, one dilatation and curettage, three vaginal hysterectomy

Follow-up:

Details are shown in table 5

Follow-up Details	
Mean follow-up±SD, range/mode (month)	6,5±5, 1-22/4
Recurrence (percentage)	6 (12,2) ²
Post operative events	9 ³
Surgery Satisfaction	33/40 ⁴
Improve in QoL	34/40 ^{4,5}

² Recurrence occur at 5,5 month average, three (6,1%) required recoloperineocleisis with optimal results in 3 month of average follow-up. ³ One De novo incontinence, one de novo constipation, seven treated with antibiotics for vaginal discharge.

⁴ Only 40 patients responded to the questionnaire. ⁵ One worse and one equal QoL.

Interpretation of results

Labhardt colpoperineocleisis is an excellent surgery for patient without sexual activity. It's fast with very low blood loss, with no intraoperative complications, few and minimal post operative complications. The rate of recurrence needing for surgery approach is very low (6,1%). This surgery improved the quality of life in this sample and it is associated with high patient's satisfaction.

Concluding message

This cohort is the biggest published data of Labhardt colpoperineocleisis, that show a fast, easy, effective and with low peri and post operative morbidity. The excellent results must be challenged in prospective studies with larger samples, so it can be installed like the gold standard in the treatment of severe genital prolapse for patients without sexual activity.

References

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Is this a clinical trial?	No
What were the subjects in the study?	HUMAN
Was this study approved by an ethics committee?	No
This study did not require ethics committee approval because	It was a retrospective cohort
Was the Declaration of Helsinki followed?	Yes
Was informed consent obtained from the patients?	Yes