Instructions to Contributors

Dear Contributor:

Enclosed in this document please find the page proofs and copyright transfer agreement (CTA) for your article in the *Journal of Coloproctology*. Please print this document and complete and return the CTA, along with corrected proofs, within 72 hours.

- Please read proofs carefully for **typographical** and **factual** errors only; mark corrections in the margins of the proofs in blue or black pen, or use **Adobe Acrobat tools** to mark the changes in the PDF file directly. Please be sure to write as clearly as possible so no errors are introduced into your article. **Answer (on the proofs) all author queries marked in the margins of the proofs.** Check references for accuracy. **Please check on the 1st page of your article that your titles and affiliations are correct.** Avoid elective changes, because these are costly and time consuming and will be made at the publisher's discretion.
- 2) Please pay particular attention to the proper placement of figures, tables, and legends. Please provide copies of any formal letters of permission that you have obtained.
- 3) Please return the corrected proofs, signed copyright transfer agreement.
- 4) After PDF approval, the author will not be able to request changes to the published article.
- 5) As a contributor to this journal you will receive a complimentary PDF file of the article after publication.

Please return all materials within 72 hours. E-mail is the easiest way to ensure your corrections are received in a timely manner. You may also return materials via fax to:

Paula Di Sessa Vavlis
Junior Project Coordinator - Journals
Thieme Revinter Publicações Ltda
Rua Rego Freitas 175, Loja 1, Bairro República
São Paulo, SP, CEP 01220-010 Brasil
Phone: +55 11 3362 2464
E-mail: paula.disessa@thieme.com.br

Please do not return your materials to the editor or the typesetter.

Please note: Due to a tight schedule, if the publisher does not receive the return of your article proofs within 7 days of the date the e-mail was sent to you, the publisher reserves the right to proceed with publication without author changes. Such proofs will be proofread by the editor and the publisher.

Thank you for your contribution to this journal.

Permission to Publish and Open Access Copyright Transfer Agreement

Manuscript Information:	
Journal:	
Manuscript Title:	
Manuscript Number:	
Authors:	
Corresponding author's contact data:	
Corresponding author's e-mail address:	
Contact at the publishers:	
E-mail address at the publishers:	

Dear Author,

Please

- read this form carefully,
- check all manuscript information,
- sign this form with your digital signature and
- reply directly to the e-mail with which you received this document and leave the subject unchanged.



Thank you very much in advance.

Assignment of Rights

Thieme does not accept for publication in a journal any manuscript that has been published elsewhere. Your consent to the following assignments of rights, also on behalf of the other authors (if several authors contribute to the manuscript), and the signing of this Open Access Copyright Transfer Agreement is a necessary requirement for the publication of your manuscript.

The article will be published under a CC BY-NC-ND license (https://creativecommons.org). This license means that anyone may freely read, download, distribute and make the article available to the public (in printed and electronic form), provided that the author and the journal as the source are acknowledged, whereas no commercial use is allowed and the work may not be altered, transformed or serve as the basis for a derivative work.

Thieme will endeavor to ensure maximum dissemination of your manuscript and to protect your authors' rights against misuse, such as plagiarism.

For this reason upon acceptance of the manuscript by us you and your co-authors agree to hereby transfer on a non-exclusive basis copyright to Thieme. This includes (in addition to the publication under a CC BY-NC-ND license), for the duration of the statutory term of copyright protection and throughout the world, the full rights to use the paper (including excerpts thereof) within the journal or separate to it, on a commercial or non-commercial basis, linked to other works or papers, audio-visual accompanying materials or interactive products or services, including the transfer to third parties (e.g., under license agreements, etc.), in particular the non-exclusive right to use all editions/updates for the following purposes:

reproduction and distribution in printed form, in particular as a journal article, article in a medical textbook or other type
of book directed towards a specific target group of readers, pocket book, special edition for secondary markets or special
customers, brochure, supplement, anthology, etc.



Permission to Publish and Open Access Copyright Transfer Agreement

- reproduction and distribution in the form of electronic media (e.g., CD-Rom, DVD, USB memory stick, databases, cloud-based service, ebook and other forms of electronic publishing) and also make available to the public (e.g., internet, intranet or other wired or wireless data networks), in particular by displaying on stationary or mobile visual display units, monitors, smart phones, tablets or other devices by download (e.g., e-pub, PDF, App) or retrieval in any other form;
- translation, transmission or adaptation of the paper into or in any other language, dialects or versions (in particular, as a podcast, audio book, or other image or sound carriers etc.), transmission through television, cable or satellite TV, radio or by means of other audio-visual media, renting or lending, storage in an electronic archive, including storage in connection with indexation, key word search or any other search and additional functions, usage for types of use not as yet known and for any other rights used by collecting societies as provided in their articles and allocation plans, provided that the transfer of such rights is permissible under the relevant provisions and applicable law.

Without prejudice to this assignment of rights, the moral rights in connection with the article shall remain with you and the other authors, whereas we are entitled to all rights with regard to the editorial process and publishing service in connection with the reproduction and distribution of the article in our journal and making it available to the public.

Thieme is entitled, but not obliged, to use the rights specified in the aforesaid sections and may adapt the article for these uses. Thieme will take your and your co-authors' legitimate interests into account in this respect.

Authors' rights and Gold Open Access

The rights of use are assigned to us non-exclusively – subject to your rights in accordance with our Open Access (OA) Guidelines: Our OA Guidelines state that immediately after the publication of the article by us, you and the other authors are entitled to make the published version of the article available to the public. For further details please click the button "Authors' rights and Gold Open Access."

For more Information on our OA Program please visit http://open.thieme.com.

Article Publication Charge

Publication of the article is subject to an article publication charge of (exclusive VAT if any).

The article will be published upon receipt of the payment by Thieme.

Duties of care

Product liability laws set high standards for your duty of care as the author of a scientific manuscript. This is especially the case when you give therapeutic information and/or specify drug doses. Therefore please check this information carefully in the typeset page-proofs of your article. Your task will be much easier if you have the information counterchecked — depending on the sensitivity of the information within the article — by specialist colleagues. Only you, as the author, have the specialist knowledge to be able to assess the accuracy of the information. For further information on how to indicate corrections, please click the button "Correction markup symbols."

Author's Declaration

I have taken note of the information on the duties of care under product liability law; I agree to the assignments of rights in accordance with the foregoing sections "Assignment of Rights" and "Authors' Rights and Gold Open Access" also on behalf of the other authors (if several authors have contributed to the article). I declare that I am authorized by my co-authors to sign on their behalf.

I declare that no third party rights will be infringed through the publication. Any material contained in the manuscript (including illustrations, tables, or other material) from third-party sources will be identified as such through citation, indicating the source. If required, I have obtained the copyright permission from the publishers concerned.

I have read and understood the terms and conditions of the CC BY-NC-ND license and I agree also on behalf of the other authors (if several authors have contributed to the article) that the article will be published under such a license.

I agree with the aforesaid article publication charge. Should one of the foregoing regulations be or become invalid in whole or in part this shall not affect the validity of the other provisions. Any invalid provision shall be replaced by a regulation that comes as close as possible to the purpose of the invalid provision in economic terms, insofar as legally permissible.

This article is ready to be published after the execution of the corrections indicated by me.

Date Digital Signature





for Authors





© Thieme Publishing Group – August 2019

Open Access Publishing – Author Information



Open Access Publishing – Author Information

Open Access (OA) means free reading and use of scientific literature and related content via the Internet.

1. OA Publishing at Thieme

There are two main ways to publish articles and make them freely available to read: "gold" OA and "green."

Gold OA means that immediately upon publication your article will be made free to read via the Internet with additional usage rights. In return for this instant and global dissemination and availability of your article, an Article Publication Charge (APC) is paid either by yourself as the author or by your institution or funding organization.

Green means that 12 months after publication you may self-archive the accepted manuscript. You may not use the final published version, but you may deposit your publication as the accepted manuscript in institutional online repositories.

Thieme offers authors the choice by providing both gold OA and green options, with the preference for gold OA as it provides immediate access to the edited, published version of your article with well-defined reuse rights under a sustainable financial model.

2. General Information — Thieme Open

Thieme is one of the market-leaders for scientific information and services in medicine and chemistry. Since the end of 2013, we have offered authors an OA publication service called Thieme Open, http://open.thieme.com.

This OA service includes rigorous peer-review, professional in-house editorial services, fast publication times, and worldwide access via Thieme's electronic journals, https://www.thieme.de/en/thieme-connect/home-3939.htm.

It also includes reference linking and long-term archiving, distribution, including the Digital Object Identifier (DOI), as well as entry into PubMed Central and indexing in OA-relevant search engines.

Thieme OA authors enjoy the following benefits – you may:

- 1. Publish under a gold OA model in our suite of gold OA journals or choose from one of our more than 150 classic journals that also offer you an OA option. Transfer your copyright and declare no conflict of interest electronically by signing the Copyright Transfer Agreement and Conflict of Interest forms.
- 2. Be certain that your OA manuscript will enjoy the same high-quality standards of peer-review and superior production services as you enjoy in all our journals.
- 3. Post the final published version of your article immediately on any site or repository if published under the gold CC BY OA option.
- 4. Deposit the final published version of your article immediately as a PDF on any non-commercial site or repository if published under the CC BY gold OA option.
- Be assured that we protect your rights as an author when you publish your article under a "CC BY-NC-ND license." Information at: https://creativecommons.org
- Be compliant with requirements, mandates, and policies of funders such as NIH, OSTP, Horizon 2020, DFG, and RCUK. For details see: http://www.sherpa.ac.uk/romeo/search.php
- 7. Be assured that your work will be archived long-term at archiving services such as CLOCKSS or Portico.
- 8. Be assured that your paper is discovered on major search engines and indexing services such as PubMed Central, Directory of Open Access Journals (DOAJ), and Scopus.



© Thieme Publishing Group – August 2019

3. Why is Open Access important for you?

OA is an additional publication service among the services and products that are provided by Thieme.

The most important reasons to publish OA are the immediate, global access, and sharing via the citable version of an article. This means a high visibility leading to a potentially higher usage, more citations, and a higher impact of your article by extended search ability and discoverability. Your article quickly gains wide scientific impact. These play an increasing role if you, your project, and the publication of your results and data are publicly funded. Please find a glossary of bibliometrics at: http://www.bibliothek.kit.edu/cms/english/bibliometrics-glossary.php

Maintaining a high-level quality standard by undergoing a rigorous peer-review process by international and leading specialists, Thieme OA articles are published quickly and may include multi-media enhancements. Furthermore, your OA publication meets relevant funder mandates, requirements, and policies.

The advantages of publishing OA in summary:



4. What does this mean for you?

What you should be considering before the transfer of your copy- and usage rights:

Your position as the copyright holder

As copyright holder you own exclusive rights on usage and re-usage of your work by third parties. The type and level of protection of copyrights and licenses vary between countries.

© Thieme Publishing Group – August 2019

The Rights Transfer - your rights as an author

When you decide to publish with Thieme, we kindly ask you to assign your copyright to us as your publisher.

OA licenses permit the copyright owner and holder to grant and impose conditions on the reuse of the content. By using a Creative Commons License you agree to a partial disclaimer of your rights as a copyright holder. You renounce irrevocably parts of your copyright and agree to the dissemination of your content to third parties free of charge.

Please find below further information about the CC licenses and their acceptance by authors:

On the CC website: https://wiki.creativecommons.org/Considerations_for_licensors_and_licensees

About the acceptance by authors: Scholarlykitchen-acceptance authors-licenses

CC BY-NC-ND license license enjoys the highest acceptance among authors, the CC BY license the lowest.

5. CC BY-NC-ND

To abide by and fully comply with our responsibility as publisher of your scientific work and record, Thieme publishes your OA paper under the most recent CC BY-NC-ND license:



CC BY-NC-ND (Attribution-Noncommercial-No Derivatives)

Users may **share**, copy, and redistribute the material in any medium or format under the following **conditions**:

Attribution — user must give appropriate credit, provide a link to the license, and indicate if changes were made. User may do so in any reasonable manner, but not in any way that suggests the licensor endorses the user or her/his

Noncommercial — user may not use the material for commercial purposes.

No Derivatives — if user remixes, transforms, or builds upon the material, the user may not distribute the modified material.

No additional restrictions — user may not apply legal terms or technological measures that legally restrict others from doing anything the license permits.

Thieme's default license is CC BY-NC-ND. However, in case of a funder requiring it, you as the author may select another license and we will arrange your publication accordingly.

Please find the relevant information on the submission page of the journal of your choice, as well as on our Copyright Transfer Agreements.

Should you have any questions or would like more information, please visit our Open Access page: http://open.thieme.com/de/web/19/faq

Or contact us: the editorial team of your journal

or

thiemeopen@thieme.de

Instructions for Author Corrections

Dear Author,

Many thanks for choosing to publish your manuscript with us! The reviewing of the typeset article by yourself is one of the last steps before publication, and we now ask for your approval to proceed with publication.

The manuscript has been prepared for publication by our copy-editors and typeset into the journal layout. Suggested changes, new formulations, deletions or restructuring of the text, if they are significant, along with questions to you are marked with a square (\blacksquare). Please consider carefully whether the changes made are suitable and are correct and respond to each question within the pdf proof. Please also check that the images have been reproduced correctly when incorporated into the layout (please note that the resolution will be higher in the print version).

During the editorial process your article has been reviewed, and (if applicable) dosages outlined in the text were scrutinized. However, only you, as the author, have the expertise to assess the accuracy and validity of the information within the text. The responsibility for the accuracy rests solely with you.

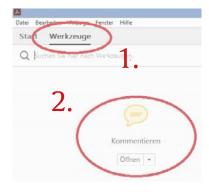
We ask that you return corrections in the following way:

To prevent misunderstandings and errors due to readabilty of handwriting, please make any necessary changes in the PDF document using the following edit functions in Adobe Acrobat <u>Reader</u> DC (freeware) or Adobe Acrobat <u>Pro</u> DC and Adobe Acrobat <u>Pro</u> (full version):

Activating the comment function

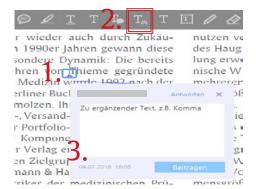
Acrobat Pro:

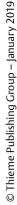
- 1. Tools
- 2. Comment



<u>Function</u>: **Inserting** text at the cursor location

- 1. Move cursor to the desired location
- 2. Click on button "Insert text at cursor"
- 3. Enter the desired text in the pop-up note box

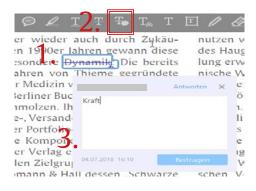




Instructions for Author Corrections

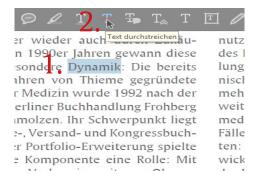
Function: Replacing text

- 1. Highlight text to be replaced
- 2. Click on button "Add note to replace text"
- 3. Enter new text in the pop-up note box



Function: Crossing out (Deleting) text

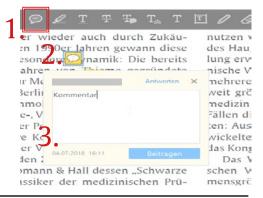
- 1. Highlight text to be deleted
- 2. Click on button "Strikethrough text"



Function: Adding a sticky note / comment

- 1. Click on button "Add sticky note"
- 2. Click on where you want to place the note in the text
- 3. Enter comments in the pop-up note box

Please never use this function for text corrections!



Please do not use the Note, Text comment and "Edit Text and Images" function as they will cause technical problems and may cause serious delay in the publication process.

You should return the PDF with the marked corrections by e-mail to your contact at the publishers.

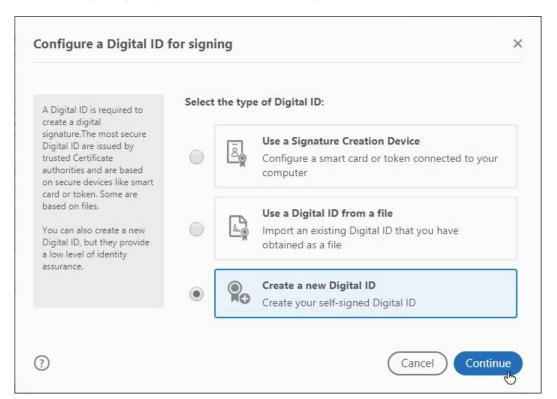
How to Set Up a Digital Signature in Adobe Reader or Adobe Acrobat

Please note: depending on the version of the Adobe program, the depiction of the dialog boxes and the wording can vary. The following screenshots were taken from Adobe Acrobat Reader DC.

To begin the digital ID setup process simply click in the appropriate signature field. Select "Configure Digital ID".

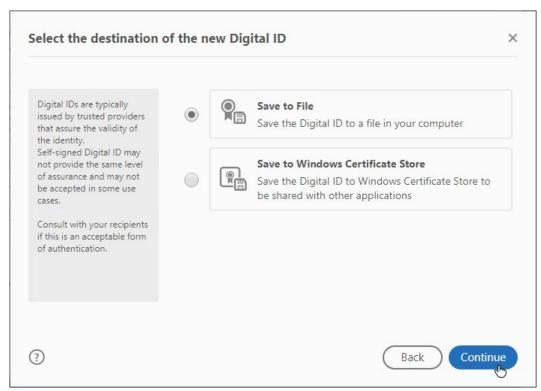


In the following dialog box, please select "Create a new Digital ID" and click Continue.

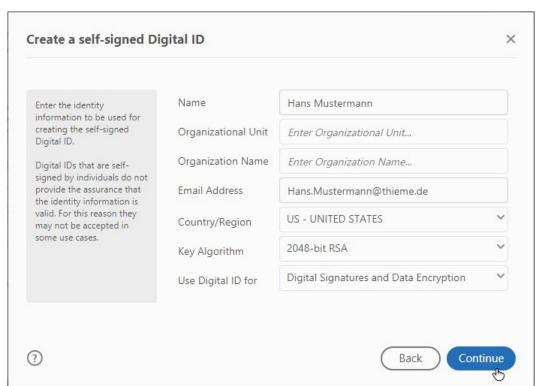




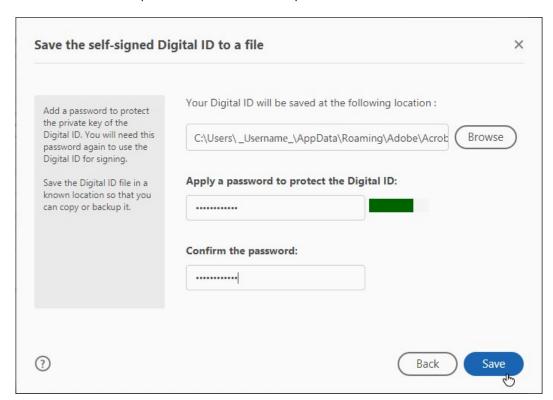
Specify where you are going to store the digital ID – select "Save to File" and click Continue.



Type in your personal information (name, organization unit, organization name and email address, country) in all fields and click Continue.



Enter a file location for your new Digital ID file – either use the default location or enter a different location if you prefer. Then create and enter a password for the ID into both password fields and click Save.







Surgical and Functional Outcome of Laparoscopic Vaginal Suspension Combined with Suture Rectopexy for Management of Pelvic Organ Prolapse

Mohamed Ibrahim Abuelnasr^{Q21} Ahmed M. F. Salama¹ Ahmed M. Nawar¹



¹Faculty of Medicine, Benha University, Egypt^{Q3}

| Coloproctol 2023;00(0):1-8.

Address for correspondence Mohamed Ibrahim Abuelnasr, M.D., Benha University Faculty of Medicine, Benha, Egypt (e-mail: drabuelnasr@gmail.com).

Abstract

Purpose Laparoscopic techniques to treat pelvic organ prolapse are gaining popularity around the globe due to their low recurrence rates and better functional results compared to perineal techniques. However, the optimum surgical procedures are not yet determined. In the current research, we suggest a novel surgical approach, laparoscopic vaginal suspension with suture rectopexy, to treat multiorgan pelvic prolapse.

Methods This prospective cohort trial was conducted from March 2018 to March 2022 and comprised 35 females with multiorgan pelvic organ prolapse with obstructed defecation symptoms. A residual rectal prolapse was still present despite the manual reduction of uterine prolapse. Patients' conditions before and after the operation were monitored regarding the obstructed defecation score, sexual function, need for laxatives, anorectal manometry pressures, anorectal sensations, and recurrence. The mean follow-up duration was one year.

Results Modified Longo score for obstructed defecation significantly decreased at six and twelve months after surgery. Additionally, a significant reduction was reported in the number of patients who needed laxatives at six and twelve months after surgery. Anorectal manometry pre- and post-surgery showed a significant elevation in the mean squeeze pressure and a decline in all rectal sensations. All parameters of the female sexual function scoring system increased postoperatively. No recurrence was reported during follow-up.

Conclusion For multiorgan pelvic prolapse, laparoscopic vaginal suspension combined with suture rectopexy has excellent functional outcomes, minimal morbidity, and low cost.

Keywords

- pelvic prolapse
- pelvic organs prolapse suspension (POPS)
- ► rectopexy

Introduction

Pelvic Organ Prolapse (POP) is a prevalent pelvic dysfunction that negatively impacts women's quality of life. It restricts their psychosocial and physical activities and affects their sexual function. POP is common in old age; it reaches 5% in 60-to 69-year-old women. It can be detected clinically in 50% of patients, with only 3% symptomatic.²

received March 11, 2023 accepted after revision October 24, 2023

DOI https://doi.org/ 10.1055/s-0043-1777082. ISSN 2237-9363.

© 2023. Sociedade Brasileira de Coloproctologia. All rights reserved.

This is an open access article published by Thieme under the terms of the Creative Commons Attribution-NonDerivative-NonCommercial-License, permitting copying and reproduction so long as the original work is given appropriate credit. Contents may not be used for commercial purposes, or adapted, remixed, transformed or built upon. (https://creativecommons.org/ licenses/bv-nc-nd/4.0/)

Thieme Revinter Publicações Ltda., Rua do Matoso 170, Rio de Janeiro, RJ, CEP 20270-135, Brazil

The POP may include the anterior and posterior vaginal walls and the vaginal apex (apical prolapses). The posterior vaginal wall prolapse is the rectal protrusion into the vagina (rectocele).³ The POP surgery objective is to rectify the prolapse and treat defecation dysfunction with simultaneous rectification of the middle-pelvic-compartment prolapse. For years, abdominal sacrocolpopexy has been preferred by specialists for treating female anterior and apical prolapse over a variety of vaginal techniques.⁴ Laparoscopic suture rectopexy may be considered an ideal laparoscopic technique for rectal prolapse, with less than a 10% recurrence rate.⁵

Longo describes pelvic organ prolapse suspension (POPS) surgery as a new technique for multi-compartment female pelvic prolapse. It is simpler than conventional techniques and significantly improves the preoperative symptoms. After POPS surgery, if a residual rectoanal prolapse or an anterior rectocele persists, a stapled transanal rectal resection (STARR) is performed.⁶

The stapled transanal rectal resection (STARR) procedure utilizes a double stapling technique with two circular staplers to remove the circumferential anorectal mucosa and fortify the anterior anorectal junction wall, rectifying the structural defects linked to ODS.⁷

The grey area about the functional outcome of a single laparoscopic technique for pelvic organ prolapse instead of POPS surgery alone has motivated the authors to conduct this study.

Patients and Methods

Study Design and Subjects:

The current prospective study was conducted between March 2018 and March 2022 at the colorectal surgery unit and the surgery department, Benha University Hospital.

It comprised 35 females with multiorgan pelvic prolapse with obstructed defecation symptoms and a residual rectal prolapse despite uterine prolapse reduction. The local ethical committee approved it (Rc5-4-2022), registered on research registry with code number (researchregistry8403) and written consent was taken from all participants.

Patients with previous surgical treatment for total rectal prolapse, history of pelvic radiotherapy, fecal incontinence, or bad general condition were excluded from the study.

Preoperative Assessment

- 1. Detailed history and Modified Longo score were used for the assessment of ODS
- 2. Clinical assessment for detection of rectal wall prolapse and its concentric folds and assessment of anal sphincter, rectocele or cystocele
- 3. Pre and postoperative anorectal manometry was done at 6 and 12 months using Solar GI High-Resolution Anal sphincter Manometry HRAM Medical Measurement Systems (MMS), Laborie, USA, with a 24- channel water perfused catheter with latex balloon to determine rectal sensations and anal pressures and exclude anismus.

Imaging

- 4. Every patient had defecography.
- 5. Colonoscopy: Each patient had a colonoscopy performed to rule out proximal lesions and get a biopsy of any rectal ulcer to rule out cancer.
- FSFI assessed sexual function before and after surgery at 6 and 12 months.
- 7. Standard preoperative laboratory examinations.

Procedure

Laparoscopic Surgical Technique

The night before the procedure, each patient underwent two rectal enemas and received 500 mg of metronidazole and 1 gm of ceftriaxone along with the induction of anaesthesia. The patient was set up in the modified lithotomy posture, with the thighs spread apart, the hips straight, and the knees flexed. Near the body were both arms.

Manual reduction of uterine prolapse and assessment of rectal prolapse were made. If residual prolapse was still present, we proceeded to our procedure. If no residual prolapse was present, we did vaginal suspension only.

The surgeon stood on the patient's right side, and the assistant and the cameraman stood on the left.

Pneumoperitoneum was produced with a Veress needle and an umbilical stab incision after urine catheterization. A 30-degree telescope was then inserted through a 12 mm trocar (camera port) that had been placed through a supraumbilical incision. The second port, which was 5 mm in size and used as the RT hand, was implanted two fingers medial to the anterior superior iliac spine. The third port, which was 5 mm in size and used as the left hand, was put at the umbilicus level at the right mid-clavicular line. The fourth port, measuring 5 mm, was placed at the left mid-clavicular line below the umbilicus for the assistant.

The patient was placed in the Trendelenburg position (30 degrees) to begin the abdominal cavity exploration. For improved anterior rectum dissection, the uterus was withdrawn to the abdominal wall using 2/0 polyproline sutures and a straight needle. The helper then pulled the sigmoid colon to the left side and out of the pelvis using the left side trocar.

Suture rectopexy was performed⁸ without preserving the lateral rectal ligaments of the lower rectum by inspecting the right ureter at the pelvic wall and performing lateral dissection through a peritoneal incision over the sacral promontory extending to the Douglas pouch.

The loose areolar tissue between the mesorectum and the presacral vein plexus was opened during the posterior rectum dissection while the presacral nerves were kept intact. Up till the pelvic floor muscles, anterior dissection was performed.

Retraction of the rectum was performed cranially to identify the ideal location for suture fixation. Before placing sutures, a rectal examination was performed to make sure there was no prolapse at the fixation point. Then, at least two interrupted polyproline 2/0 sutures were used to attach the seromuscular layer of the rectum posterior wall to the presacral fascia on both sides (**Fig. 1**).



Fig. 1 (A) Posterior dissection of the rectum. (B) Anterior dissection of the rectum. (C) Suturing rectum to presacral fascia.

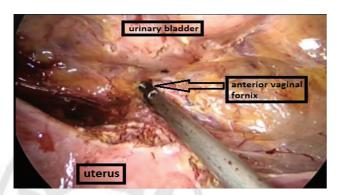


Fig. 2 Dissection done till reaching anterior vaginal fornix.

For greater suspension and to prevent adhesions, the lateral peritoneum was continuously sutured to the rectum at the new higher location. The sutures used were PDS 2/0.

The vaginal suspension described by Longo⁹ was done by cutting the sutures used to retract the uterus to the abdominal wall. A peritoneal incision (3 cm) between the uterus and urinary bladder was done. Then, the dissection was started at the anterior vaginal fornix apex (►Fig. 2).

A V-shaped mesh (25 cm long, 2 cm wide) was prepared and introduced through a 10 mm trocar into the abdominal cavity. The mesh apex was fixed using a PDS 2/0 by three sutures on the anterior vaginal vault.

A subperitoneal tunnel was done on both sides by the Endo Clinch to reach the anterior vaginal fornix. The subperitoneal tunnel was used to extract each end of the V-mesh, and symmetrical traction was applied to both

mesh strips. (Fig. 3) When the vaginal vault was suspended at the ideal level to eliminate the vaginal prolapse while preventing excessive tension on the vaginal walls, the second assistant informed the surgeon. The correction happened following CO2 exsufflation. Each mesh strip was fixed to the external oblique aponeurosis by prolene 0 stitches.

After reinsufflaton, the peritoneum was closed by continuous sutures using vicryl 2/0.

For all included patients crucial monitoring intraoperative and post operative for

- Intraoperative data, including blood loss, operative time, and complications.
- Postoperative data, including pain score, hospital stay, complications, and the time needed for mesh incorporation.
- Follow-up was extended for 12 months for the incidence of recurrence of ODS and
- Obstructed defecation symptoms by the modified Longo score. 10
- Anorectal manometry and rectal sensations.
- The need for laxatives postoperatively.
- Sexual function by the Female Sexual Function Index (FSFI), which is a 19-item It comprises six domains: desire [two items], lubrication [four items], arousal [four items], pain, orgasm, and satisfaction [three items each]. 11
- The FSFI total score is the sum of the six domain/subscale ratings, and the maximum score is 36. Higher scores imply greater functioning, and a total score of 26 is the diagnostic threshold for female sexual dysfunction.

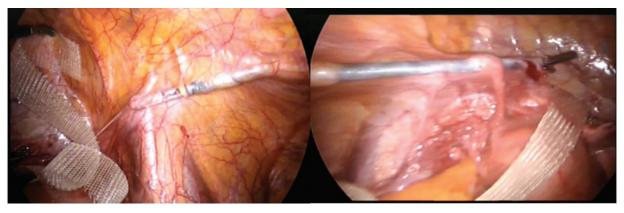


Fig. 3 Creation of right and left subperitoneal tunnel and pulling the right limb of mesh.

 All functional outcomes were assessed at six and twelve months postoperatively and compared to the preoperative ones.

Postoperative Care and Instructions

Ceftriaxone (1 gm) was given once /day for five days. Analgesia was given as patient-controlled analgesia (PCA) pump. Early postoperative laxatives were prescribed to avoid straining and discontinued gradually.

The visual analogue score (VAS) was used to assess postoperative pain. Patients started taking oral fluids on the surgery day evening and soft food the day after surgery.

All patients were admitted on the surgery day. The hospital stay duration was calculated from admission to the day of discharge.

Outcomes and Follow Up

Primary outcome: The 1ry outcome was successful POPS surgery with relief of obstructed defecation symptoms.

Secondary outcomes: The 2 ry outcome was improvement of the sexual function QOL score.

Postoperative follow-up was done in the outpatient clinic one week after the operation and then every month for 12 months.

At 6 and 12 months, patients were reassessed by anorectal manometry and modified Longo score. Additionally, sexual function was assessed by FSFI. In the outpatient clinic, the questionnaire was distributed by a nurse and administered by the patients. Recurrence was evaluated clinically and monitored for one year or until the study ended.

Statistical Analysis

The primary goal of this study is to reduce the occurrence of ODS symptoms; hence the sample size was estimated accordingly. Using the G*power 3.1 programme (Universities, Dusseldorf, Germany), a sample size of 35 was considered with a power of 80%, P value of 0.05, and an effect size of 0.7.

By examining the data distribution, the normality of the measured results was investigated. A parametric distribution was visible for all examined parameters. The IBM Corp., Armonk, NY, USA used a two-way mixed model MANOVA to compare measured variables over several time periods. Independent t test was used for participant demographic information, and 2 was used for nominal information. Nominal data was given as a number and a percentage whereas numerical data was presented as mean and SD. With a P value of less than or equal to 0.05, the significance level was established. The statistical analysis was performed using SPSS Statistics version 20.

Results

This study included 35 females with multiorgan pelvic prolapse with obstructed defecation. A residual rectal prolapse was still present despite the manual reduction of uterine prolapse during examination under general anesthesia.

Table 1 Baseline characteristics in the study participants

	n =35
Age (years) Gender (Female) BMI (kg/m²) Comorbidities Main complaint	33 ± 8.12 35 (100%) 39.03 ± 11.81 9 (25.7%)
OBD OBD + Sexual complaint	26 (74.3%) 9 (25.7%)

Data are presented as mean \pm standard deviation (SD) or n (%), BMI: Body mass index, OBD: Obstructive bowel disorder.

As shown in **-Table 1**, the mean age of the studied patients was 33 ± 8.12 years. The mean BMI was 39.03 ± 11.81 . Nine patients (25.7%) had comorbidities. The main complaint was OBD in 26 patients (74.35%), while OBD with sexual complaints was reported in 9 (25.7%).

► **Table 2** shows intra and postoperative findings. The ASA score was 1 in 12 patients (34.3%), 2 in 19 (54.3%), and 3 in only 4 (11.4%). The median operative time was 90 minutes. The median blood loss was 20 ml. No cases were converted to open. Concerning the intraoperative complications, only two patients (5.71%) had bleeding. No intestinal, uterine, or bladder injuries occurred. Regarding postoperative findings, the median pain score was 4. Wound infection occurred in two patients (5.7%). Only one patient (2.9%) had a hematoma. No recurrence was reported. The median hospital stay was 2.2 days.

►Table 3 and **►Figure 4** show the change in anorectal manometry parameters at 6 and 12 months postoperatively. Compared to baseline values, the mean resting and squeezing pressures significantly increased at 6 and 12 months postoperatively (P < 0.001 for each). In contrast, 1st rectal sensation volume significantly decreased at 6 and 12 months postoperatively (50 and 60 cc, respectively, vs. 70 cc,

Table 2 Intra and postoperative data in the study participants

	n =35
*Operative data	
ASA score	12 (34.3%)
1	19 (54.3%)
2	4 (11.4%)
3	90 (90 - 100)
Operative time (minutes)	20 (20 - 30)
Blood loss (mL)	0 (0%)
Conversion to open	
*Intraoperative complications	
Bleeding	2 (5.71%)
Intestinal injury	0 (0%)
Uterine or bladder injury	0 (0%)
*Post operative data	
Post operative pain score	4 (2 - 4)
Wound infection	2 (5.7%)
Hematoma	1 (2.9%)
Hospital stay (days)	2.2 (2 - 3)
Recurrence	0 (0%)

Data are presented as mean \pm standard deviation (SD), median (IQR), or n (%), ASA: American Society of Anesthesiologists

Table 3 Anorectal manometry preoperatively and in 6 and 12 months postoperatively

	Preoperative	6 months postoperative	12 months postoperative	P value
Mean resting pressure	35 (30 - 75) ^a	40 (35 - 50) ^b	40 (40 - 50) ^b	< 0.001
Mean squeezing pressure	110 (100 - 120) ^a	130 (120 - 160) ^b	140 (130 - 160) ^b	<0.001
1 st rectal sensation volume (cc)	70 (60 - 80) ^a	50 (40 - 60) ^b	60 (50 - 60) ^b	<0.001
1 st urge sensation volume (cc)	140 (130 - 170) ^a	130 (130 - 150) ^a	150 (130 - 160) ^a	0.034
Intense urge sensation volume (cc)	230 (210 - 260) ^a	210 (190 - 230) ^b	220 (200 - 230) ^b	<0.001

Data are presented as mean \pm standard deviation (SD), median (IQR), or n (%). Different super-scripted letters indicate significant difference and vice versa. Level if significance were set at p value \leq 0.05.

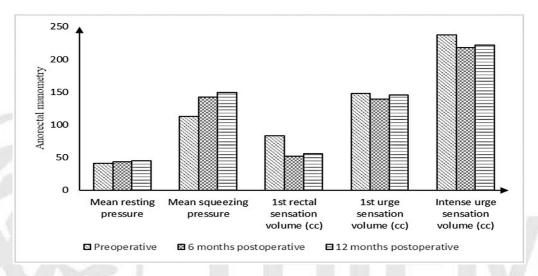


Fig. 4 Anorectal manometry preoperatively and in 6 and 12 months postoperatively.

P < 0.001). Additionally, intense urge sensation volume significantly decreased at 6 and 12 months postoperatively (210 and 220 cc, respectively, vs. 230 cc, P < 0.001). No significant differences were reported between the six- and twelvemonth values of resting and mean squeezing pressures, 1st rectal sensation volume, and intense urge sensation volume. Furthermore, pre and postoperative 1st urge sensation volume did not significantly differ.

As shown in **►Table 4**, the modified Longo score significantly declined at 6 and 12 months postoperatively compared to baseline (10 and 8, respectively, vs. 18, P < 0.001), indicating an improvement in ODS. In contrast, no significant difference was observed between the six- and twelve-month scores. A lower number of patients needed laxatives at 6 and 12 months postoperatively (6 and 5 patients, respectively) compared to baseline (35) (P < 0.001).

► **Table 5** and ► **Figure 5** reveal the change in FSFI score at 6 and 12 months postoperatively. Desire, arousal, and orgasm

scores significantly increased at 6 months compared to baseline. In addition, they significantly increased at 12 months compared to 6 months and baseline (P < 0.001 each). Lubrication, satisfaction, and pain scores significantly increased at 6 months compared to baseline, but no significant differences were observed between six and twelve months. The total FSFI score significantly increased at six months (21.34 \pm 1.68) compared to baseline (14.03 \pm 1.62). Additionally, it significantly increased at 12 months (25.32 ± 2.2) compared to 6 months and baseline.

Discussion

POP is a disability that typically affects older women and alters their lifestyles. Over 23% of community-dwelling women will have a minimum of one of the following, obstructed defecation syndrome (ODS), pelvic organ prolapses (POP), or fecal incontinence. 12 Half of the women

Table 4 Modified Q4Longo score and need for laxatives preoperatively and in 6 and 12 months postoperatively

	Preoperative	6 months postoperative	12 months postoperative	P value
Modified longo score	18 (16–20) ^a	10 (8–12) ^b	8 (8–10) ^b	<0.001
Need for laxatives	35 (100%) ^a	6 (17.14%) ^b	5 (14.28%) ^b	<0.001

Data are presented as mean \pm SD or median (IQR), or n (%). Different super-scripted letters indicate significant difference and vice versa. FSFI: Female Sexual Function Index. Level of significance were set at p value ≤ 0.05 .



Table 5 FSFI score preoperatively and in 6 and 12 months postoperatively

	Preoperative	6 months postoperative	12 months postoperative	P value
Desire	2.4 (1.2-2.4) ^a	3.6 (3.6–3.6) ^b	4.8 (3.6–4.8) ^c	< 0.001
Arousal	2.1 (2.1–2.4) ^a	3 (2.4–3.6) ^b	3.6 (3–3.9)°	< 0.001
Lubrication	$2.4(2.4 - 3)^{a}$	3.6 (3–3.9) ^b	3.9 (3.6–4.2) ^b	< 0.001
Orgasm	2.8 (2.4–3.2) ^a	$3.6 (3.2 - 4)^a$	4.8 (4–4.8) ^c	< 0.001
Satisfaction	2 (1.6–2.4) ^a	4 (3.2–4.8) ^b	4.8 (4–5.6) ^b	< 0.001
Pain	2 (1.6–2.8) ^a	4 (3.2–4.8) ^b	4.4 (4–5.2) ^b	< 0.001
Total score	14.03 ± 1.62^{a}	21.34 ± 1.68 ^b	25.32 ± 2.2°	< 0.001

Data are presented as mean \pm SD, median (IQR), or n (%). Different super-scripted letters indicate significant difference and vice versa. FSFI: Female Sexual Function Index. Level if significance were set at p value \leq 0.05.

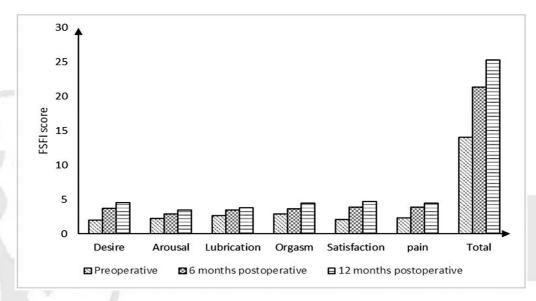


Fig. 5 FSFI score preoperatively and in 6 and 12 months postoperatively.

with ODS caused by rectal prolapse and rectocele also had uterine prolapse.¹³ Thirty percent of women who underwent POP surgery require further pelvic surgery.¹⁴

In 2014, Longo et al. decided to improve pelvic organ prolapse surgery to achieve simultaneous prolapse correction of all pelvic organs and remission of the associated symptoms by POPS surgery, which is considered a one-stage treatment for multiorgan female pelvic prolapse. Ceci et al. state that laparoscopic POPS has outstanding morphological outcomes for vaginal prolapse, enterocele, rectocele, and rectal prolapse, with no complications or functional impairments as dyspareunia or mesh-related erosion of the rectum. Nevertheless, it can be linked to a persistent and recurrent rectal prolapse. Therefore, they concluded that the STARR technique should be utilized if a residual rectal prolapse continues.⁶

The STARR technique utilizes a double stapling approach with two circular staplers to remove the circumferential anorectal mucosa and fortify the anterior anorectal junction wall, rectifying the structural defects linked to ODS. On the other hand, using two circular staplers, which cost about 1250 USD, is considered very expensive in such a low-income country and is not covered by the insurance company in most cases. In addition, the time lost in changing patients' positions from being supine during the POPS procedure to a

lithotomy position in the STARR procedure in the operative room makes the procedure more time-consuming.

Based on this background, for patients with multiorgan prolapse in which residual rectal prolapse is still present after vaginal suspension, we modified the POPS procedure and proposed a new combined laparoscopic technique, laparoscopic vaginal suspension and rectopexy, in which middle and posterior pelvic compartment prolapse can be corrected.

We preferred adding suture rectopexy to vaginal suspension over ventral mesh rectopexy for two reasons. First, to avoid many mesh-related complications such as pain, dyspareunia, rectovaginal fistula, and rectal stricture. Second, vaginal suspension alone corrects the rectocele by stretching the posterior vaginal wall but does not optimally correct rectoanal intussusception, particularly if the intussusception is of a high level, necessitating suture rectopexy rather than ventral mesh rectopexy. In 2017, Yang et al. modified POPS surgery for rectal and uterine prolapse by adding suture rectopexy to vaginal suspension using a trimmed 3-strip mesh. ¹⁵

In this study, obstructed defecation symptoms were observed in 35 females with multiorgan pelvic organ prolapse. These symptoms were assessed preoperatively by the modified Longo score. Postoperatively, significant improvements

in ODS score were observed, which may be due to the surgical approach utilized. The lateral ligaments were separated for complete rectal mobilization, and the posterior vaginal wall was stretched by vaginal suspension to correct the rectocele. Finally, the lateral peritoneum was sutured to the rectum at a new higher point for more suspension. Despite improvement in ODS, five patients (14%) depended on laxatives, which may be due to these patients' slow colonic transit time, so we recommend resection rectopexy to those patients despite the risk associated with colonic anastomosis.¹⁶

Women with pelvic organ prolapse experience impairments in sexual function. In the current study, the sexual activity of all patients was assessed pre and postoperative using the Female Sexual Function Index Scoring. Nine patients (26%) preoperatively complained of sexual dysfunction, while postoperatively, all parameters, including arousal, desire, orgasm, satisfaction, lubrication, and pain scores, significantly improved. However, only one patient had a worsened pain score postoperatively, which may be a reaction to the mesh inserted in the anterior vaginal fornix. She was managed conservatively with non-steroidal analgesics, and the pain gradually decreased with time. The 26 patients (74%) who did not complain of preoperative sexual problems reported better sexual function postoperatively, and all parameters were significantly increased.

Treating genital prolapse has a good impact on sexual activity. It helps eliminate vaginal dryness and chronic pelvic pain related to uterine prolapse and symmetrical traction of the mesh and avoids excessive shortening of the vagina. Rogers reports that surgical treatment of the prolapse improves sexual function and body image. This has been demonstrated in both native tissue and grafted repairs in most patients.¹⁷ However, a meta-analysis by Abed et al. states that roughly 10% of mesh-repaired prolapse patients experience dyspareunia after surgery.¹⁸

The recurrence incidence is a key metric for evaluating pelvic prolapse surgery. In this study, no recurrence was reported postoperatively, which could be attributed to adding suture rectopexy to vaginal suspension, full mobilization of the rectum to the pelvic floor, and bilateral suturing of the rectum to the sacral promontory. While in a study done by Yang et al. in which modification POPS surgery were done using a trimmed 3-strip mesh, 1 patient had a recurrence from 69 patients included in the study.¹⁵

The current study had limitations, such as the small sample size and the limited follow-up. In addition, it is a single-center study. We also did not spot the light on urological problems accompanied by pelvic organ prolapse.

Conclusion

The laparoscopic vaginal suspension combined with suture rectopexy has outstanding functional outcomes, minimal morbidity, and a low cost compared to POPS described by Longo (when STAR operation is needed). In addition, it is associated with a short hospital stay, alleviation of obstructed defecation symptoms, low recurrence rate, minimal mesh-related complications and postoperative functional impairment, and correction of middle-compartment prolapses. The long-term outcomes will be achieved by follow-up.

Conflicts of Interest

Acknowledgment

None

References

- 1 Belayneh T, Gebeyehu A, Adefris M, Rortveit G, Awoke T. Pelvic organ prolapse in Northwest Ethiopia: a population-based study. Int Urogynecol J Pelvic Floor Dysfunct 2020;31(09):1873–1881
- 2 Wu JM, Vaughan CP, Goode PS, et al. Prevalence and trends of symptomatic pelvic floor disorders in U.S. women. Obstet Gynecol 2014;123(01):141-148
- 3 Iglesia CB, Smithling KR. Pelvic Organ Prolapse. Am Fam Physician 2017;96(03):179-185
- 4 Maher C, Feiner B, Baessler K, Schmid C. Surgical management of pelvic organ prolapse in women. Cochrane Database Syst Rev 2013;(04):CD004014
- 5 Tou S, Brown SR, Nelson RL. Surgery for complete (full-thickness) rectal prolapse in adults. Cochrane Database Syst Rev 2015;2015 (11):CD001758
- 6 Ceci F, Spaziani E, Corelli S, et al. Technique and outcomes about a new laparoscopic procedure: the Pelvic Organ Prolapse Suspension (POPS). G Chir 2013;34(5-6):141-144
- Altomare DF, Rinaldi M, Veglia A, Petrolino M, De Fazio M, Sallustio P. Combined perineal and endorectal repair of rectocele by circular stapler: a novel surgical technique. Dis Colon Rectum 2002;45(11):1549-1552
- 8 Dulucq JL, Wintringer P, Mahajna A. Clinical and functional outcome of laparoscopic posterior rectopexy (Wells) for fullthickness rectal prolapse. A prospective study. Surg Endosc 2007;21(12):2226-2230
- 9 Longo A, Boller B, Crafa F, Perrone F. Pelvic organ prolapse suspension. Pelvic Floor Disorders: Surgical Approach. 2014: 207-217
- 10 Sharma S, Agarwal B. Scoring systems in evaluation of constipation and obstructed defecation syndrome (ODS). JIMSA 2012;25 (01):57-59
- 11 Sand M, Rosen R, Meston C, Brotto L. The female sexual function index (FSFI): a potential "gold standard" measure for assessing therapeutically-induced change in female sexual function. Fertil Steril 2009;92(03):S129
- 12 Nygaard I, Barber MD, Burgio KL, et al; Pelvic Floor Disorders Network. Prevalence of symptomatic pelvic floor disorders in US women. JAMA 2008;300(11):1311-1316
- 13 González-Argenté FX, Jain A, Nogueras JJ, Davila GW, Weiss EG, Wexner SD. Prevalence and severity of urinary incontinence and pelvic genital prolapse in females with anal incontinence or rectal prolapse. Dis Colon Rectum 2001;44(07):920-926
- 14 Olsen AL, Smith VJ, Bergstrom JO, Colling JC, Clark AL. Epidemiology of surgically managed pelvic organ prolapse and urinary incontinence. Obstet Gynecol 1997;89(04):501-506
- 15 Yang SJ, Yoon SG, Lim KY, Lee JK. Laparoscopic Vaginal Suspension and Rectopexy for Rectal Prolapse. Ann Coloproctol 2017;33(02):64-69

- 16 Formijne Jonkers HA, Maya A, Draaisma WA, et al. Laparoscopic resection rectopexy versus laparoscopic ventral rectopexy for complete rectal prolapse. Tech Coloproctol 2014;18(07): 641–646
- 17 Rogers RG. Sexual function in women with pelvic floor disorders. Can Urol Assoc J 2013;7(9-10, Suppl 4)S199–S201
- 18 Abed H, Rahn DD, Lowenstein L, Balk EM, Clemons JL, Rogers RGSystematic Review Group of the Society of Gynecologic Surgeons. Incidence and management of graft erosion, wound granulation, and dyspareunia following vaginal prolapse repair with graft materials: a systematic review. Int Urogynecol J Pelvic Floor Dysfunct 2011;22(07):789–798



Author Query Form (JCOL/2300030)

Special Instructions: Author please write responses to queries directly on proofs and then return back.

- <u>O1</u>: AU: Please provide a short version of the title.
- Q2: AU: Please confirm that given names (red), middle names (black) and surnames (green) have been identified correctly. Author names in bibliographic citations and online will appear as: Abuelnasr MI, Salama AMF, Nawar AM. Surgical and Functional Outcome of Laparoscopic Vaginal Suspension Combined with Suture Rectopexy for Management of Pelvic Organ Prolapse. Please confirm if this is correct.
- <u>Q3</u>: AU: Please provide the department of this affiliation.
- <u>O4</u>: AU: Please provide significance of the tables 3, 4 and 5.

