

U. PORTO

FMUP FACULDADE DE MEDICINA
UNIVERSIDADE DO PORTO

MESTRADO INTEGRADO EM MEDICINA

2018/2019

Andreia Vanessa Almeida Pisco

Abdominoplastia com Preservação da Fáscia de Scarpa:

Estudo Prospetivo Comparativo sobre Número de Drenos/

Abdominoplasty with Scarpa Fascia Preservation:

Prospective Comparative Study of Suction Drain Number

março, 2019

FMUP

U. PORTO

FMUP FACULDADE DE MEDICINA
UNIVERSIDADE DO PORTO

Andreia Vanessa Almeida Pisco

Abdominoplastia com Preservação da Fáscia de Scarpa:

Estudo Prospetivo Comparativo sobre Número de Drenos /

Abdominoplasty with Scarpa Fascia Preservation:

Prospective Comparative Study of Suction Drain Number

Mestrado Integrado em Medicina

Área: Cirurgia Plástica, Reconstructiva e Estética

Tipologia: Dissertação

Trabalho efetuado sob a Orientação de:

Professor Doutor António Costa-Ferreira

E sob a Coorientação de:

Professora Doutora Helena Peres

Trabalho organizado de acordo com as normas da revista:

Annals of Plastic Surgery

março, 2019

FMUP

Eu, Andreia Vanessa Almeida Pisco, abaixo assinado, nº mecanográfico 201202276, estudante do 6º ano do Ciclo de Estudos Integrado em Medicina, na Faculdade de Medicina da Universidade do Porto, declaro ter atuado com absoluta integridade na elaboração deste projeto de opção.

Neste sentido, confirmo que **NÃO** incorri em plágio (ato pelo qual um indivíduo, mesmo por omissão, assume a autoria de um determinado trabalho intelectual, ou partes dele). Mais declaro que todas as frases que retirei de trabalhos anteriores pertencentes a outros autores, foram referenciadas, ou redigidas com novas palavras, tendo colocado, neste caso, a citação da fonte bibliográfica.

Faculdade de Medicina da Universidade do Porto, 22/3/2019

Assinatura conforme cartão de identificação:

Andreia Vanessa Almeida Pisco

NOME

Andreia Vanessa Almeida Pisco

NÚMERO DE ESTUDANTE

E-MAIL

201202276

avalemeidap@hotmail.com

DESIGNAÇÃO DA ÁREA DO PROJECTO

Cirurgia Plástica, Reconstructiva e Estética

TÍTULO DISSERTAÇÃO/~~MONOGRAFIA~~ (riscar o que não interessa)

Abdominoplasty with Scarpa Fascia Preservation: Prospective Comparative Study of Suction Drain Number

ORIENTADOR

António Manuel Domingues da Costa Ferreira

COORDENADOR (se aplicável)

Maria Helena Tabuaço Rego Martins Peres

ASSINALE APENAS UMA DAS OPÇÕES:

É AUTORIZADA A REPRODUÇÃO INTEGRAL DESTES TRABALHOS APENAS PARA EFEITOS DE INVESTIGAÇÃO, MEDIANTE DECLARAÇÃO ESCRITA DO INTERESSADO, QUE A TAL SE COMPROMETE.	<input checked="" type="checkbox"/>
É AUTORIZADA A REPRODUÇÃO PARCIAL DESTES TRABALHOS (INDICAR, CASO TAL SEJA NECESSÁRIO, Nº MÁXIMO DE PÁGINAS, ILUSTRAÇÕES, GRÁFICOS, ETC.) APENAS PARA EFEITOS DE INVESTIGAÇÃO, MEDIANTE DECLARAÇÃO ESCRITA DO INTERESSADO, QUE A TAL SE COMPROMETE.	<input type="checkbox"/>
DE ACORDO COM A LEGISLAÇÃO EM VIGOR, (INDICAR, CASO TAL SEJA NECESSÁRIO, Nº MÁXIMO DE PÁGINAS, ILUSTRAÇÕES, GRÁFICOS, ETC.) NÃO É PERMITIDA A REPRODUÇÃO DE QUALQUER PARTE DESTES TRABALHOS.	<input type="checkbox"/>

Faculdade de Medicina da Universidade do Porto, 22/3/2019

Assinatura conforme cartão de identificação: Andreia Vanessa Almeida Pisco

Ao meu avô, que me ensinou que a curiosidade anda a par com a felicidade, e que me mostrou que o conhecimento é a porta para o mundo. Por sempre ter acreditado!

À minha avó, que me ensinou a contar, ler e escrever, mas mais importante ainda, a viver a vida com um sorriso nos lábios.

Aos meus pais, que estão sempre lá. Pela confiança, pela paciência, por compreenderem as ausências. Por darem tudo de si para eu ser o que sou hoje, sem pedir nada em troca.

À minha irmã, que consegue sempre pôr-me um sorriso nos lábios. Por me ouvir de forma incondicional. Pelas noitadas no sofá.

Às minhas amigas, pelo encorajamento ilimitável, pelas gargalhadas capazes de afastar qualquer nuvem negra e por celebrarem comigo cada passo desta longa caminhada.

Ao Professor António Costa-Ferreira, por todo o apoio e atenção durante a elaboração deste trabalho e pelo conhecimento transmitido, que levarei comigo para a minha carreira futura.

Abdominoplasty with Scarpa Fascia Preservation: Prospective Comparative Study of Suction Drain Number

Authors:

Andreia Vanessa Almeida Pisco, Medical Student *

Maria Helena Tabuaço Rego Martins Peres, Ph.D. †

António Manuel Domingues da Costa-Ferreira, M.D., Ph.D. ‡

Affiliation:

* Porto University Medical School, Porto, Portugal,

† Interdisciplinary Centre of Marine and Environmental Research (CIIMAR) and Science Faculty, Porto University, Porto, Portugal

‡ Department of Plastic, Reconstructive and Aesthetic Surgery and Burn Unit, São João Hospital, Porto University Medical School, Porto, Portugal

Correspondence and Reprints: António Costa-Ferreira, M.D., Ph.D., Department of Plastic, Reconstructive and Aesthetic Surgery and Burn Unit, São João Hospital, Porto University Medical School, Alameda Prof. Hernâni Monteiro, 4200-319 Porto, Portugal.
Contact: 220426676 E-mail: antoniof@med.up.pt

Conflicts of Interest and Source of Funding: none declared.

ABSTRACT

BACKGROUND: Abdominoplasty is becoming increasingly more common, with seroma being the most frequent complication. Suction drains are used very often as a method to prevent seroma formation and it has been suggested that techniques using Scarpa fascia preservation and closed-suction drains have lower seroma rates than other approaches. However few studies have addressed parameters that may affect drain efficiency. A prospective comparative study was conducted to determine if applying two or three closed-suction drains, after an abdominoplasty with Scarpa fascia preservation, has any effect on several outcomes.

METHODS: This was a single-center study conducted from September 2016 to March 2019. Patients were allocated according to choice to one of the two surgeons involved in the study, each responsible for one group: abdominoplasty with Scarpa fascia preservation with two closed-suction drains placed postoperatively (group A) or with three closed-suction drains (group B). A comparative analysis of selected variables was done between both groups, including time to drain removal, total and daily drain output, duration of hospital stay, emergency department visit, readmission to the hospital, secondary surgical procedure, and incidence of postoperative local and systemic complications.

RESULTS: A total of 73 abdominoplasties with Scarpa fascia preservation were performed in women (group A, 33 patients; group B, 40 patients). General characteristics of group A and B were similar. There were no statistically significant differences between groups in any of the determined variables, namely main outcomes (total and daily drain output, time to drain removal) or complications (local or systemic).

CONCLUSIONS: Our results suggest that using three closed-suction drains postabdominoplasty with Scarpa fascia preservation has no advantages in total and daily drain output, time to drain removal or complications when compared with the usual two drains approach.

Level of Evidence: Level II

KEY WORDS: abdominoplasty, Scarpa fascia preservation, suction drain, drain number, drain efficiency

INTRODUCTION

Abdominoplasty is amongst the most common cosmetic surgical procedures, and like the majority of surgical interventions, it has seen several innovations in the past few decades, mainly in order to reduce complications and improve patient satisfaction.¹⁻⁶ Being seroma the most frequent complication associated with abdominoplasty, with an incidence ranging from 5 to 50%,^{4, 6-14} this is often the focus when evaluating new techniques and approaches.

Suction drains are frequently used postabdominoplasty, as a method to prevent seroma formation.^{4, 13, 15-17} The widest survey on abdominoplasty, by Matarasso et al., reported that 98% of inquired surgeons used suction drains on their approach.¹⁵ In the largest clinical series published, with 1008 abdominoplasties over an 11 years period, all 6 surgeons involved used suction drains.⁴

It has been shown that using abdominoplasty with Scarpa fascia preservation has several important advantages, as it significantly reduces drain output and time to drain removal, as well as seroma rates,^{6, 18-22} while also avoiding long drainers completely and providing identical aesthetic results.^{6, 22} Quaba et al. reported that techniques using Scarpa fascia preservation and closed-suction drains in the postoperative period have the lowest seroma rates in the literature (0 to 2.5%), when compared with studies using traditional abdominoplasty with drains (5 to 19%), progressive tension sutures without use of drains (0.2 to 8.8%) or Scarpa fascia preservation and no use of drains (7.7%).⁵ Two clinical series analysing techniques using Scarpa fascia preservation and no-drains have both failed in accomplishing seroma rates as low as the ones registered with the use of Scarpa sparing techniques with drains.^{5, 23} Several studies have shown that using

progressive tension “quilting” sutures significantly reduces the rates of seroma^{2, 14, 24} and renders drains unnecessary,^{1, 3} but this method requires additional training, increases intraoperative time and does not achieve seroma rates as low as the ones obtained with Scarpa sparing and drains.^{3, 13, 24}

The fact is that closed suction drains are still widely used and should be considered whenever dead space is created.^{16, 17} However, few studies have addressed parameters that may affect drain efficiency. There are no studies comparing the intensity of drain vacuum in abdominal procedures. Due to lack of studies, the optimal parameters for closed-suction drain use are not well known and its use and care, by surgeons, nurses, and caregivers, may be inconsistent.

To the best of our knowledge, there are no studies on the number of drains and on what might be the best approach to use in abdominoplasty. Our study aims to clarify this matter with a prospective comparative study to determine if applying two or three closed-suction drains, after an abdominoplasty with Scarpa fascia preservation, has any effect on several outcomes.

PATIENTS AND METHODS

This was a prospective comparative study, at a single institution, conducted in Porto, Portugal, at Hospital of Santa Casa da Misericórdia de Lousada, from September 2016 to March 2019. Approval was granted by the Ethical Committee of this institution and Informed Consent was given by all involved patients.

Seventy-three consecutive patients who sought treatment with the involved surgeons were selected. Eligibility criteria were: female patients who presented

abdominal deformity, marked by excess abdominal skin and adipose tissue with muscle laxity, and met the criteria for a full abdominoplasty with umbilical transposition (Psillakis types III and IV and Matarasso types III and IV).^{25, 26} Exclusion criteria were: significantly elevated operative health risks, bariatric patients without weight stabilization for at least 6 months, patients who anticipate future pregnancy, and patients with a body mass index over 30 kg/m², except those with previous bariatric surgery. Active smokers were instructed to stop smoking or to reduce smoking to 3 cigarettes per day 6 weeks before surgery, being considered as active smokers.

Two fully trained surgeons were involved in the study. Patients were allocated to each surgeon according to patient choice, but both surgeons worked together during all surgeries, differing on who was the main surgeon. Surgeon A performed an abdominoplasty with Scarpa fascia preservation and using two suction drains postoperatively (Group A), while Surgeon B performed the exact same surgical procedure with the sole difference of using three suction drains (Group B). The anatomical areas where drains were placed in each group are shown in Figure 1.

Data were extracted from patients' clinical charts, which included general demographics and clinical characteristics, intraoperative details, postoperative regimen, complications, and follow-up.

Surgical Methods

All patients routinely received preoperative enoxaparin (40mg/day subcutaneously during the hospital stay starting at least 2 hours before surgery) and broad-spectrum intravenous antibiotics. The surgical procedure began with preparing and draping the patient under general anaesthesia. All patients were submitted to a full

abdominoplasty with umbilical transposition, rectus abdominis plication and preservation of the Scarpa fascia and the deep fat compartment in the infraumbilical area (the abdominal flap was dissected by electrocautery in two different planes: pre-Scarpa fascia in the lower abdomen and pre-muscular in the epigastric region and infraumbilical midline). Both the preoperative markings and the surgical technique of this procedure are well described elsewhere.^{6, 20, 22}

The diathermocoagulation device used was ERBE VIO 300 S (Erbe Elektromedizin GmbH; Tuebingen, Germany), set to program 6, effect 4, with coagulation and cut regulated to 60, spray mode deactivated. The settings were always the same for both surgeons and for all the procedures.

Liposuction was limited to the flanks and no additional procedures were performed in the same operative time. No quilting sutures were used.

Two closed-suction drains were used in group A, placed in the right and left iliac fossae and in group B an additional third closed-suction drain was placed extending to the epigastric area (Figure 1). Drains used were VyDrain 600ml, with an initial vacuum pressure of 900 mbar (Vygon; Ecoen, France), together with a Redon drainage tube of 50 cm, 15 cm of which are double perforated (Braun; Melsungen, Germany).

The procedure did not differ in any other aspects between both groups. Compression garments were routinely used and applied in the operating room and the patients were motivated to ambulate on the first postoperative day. Drains were routinely removed when the patient was ambulatory and the single drain output was equal or less than 50 ml, collected over 24 hours, but were never removed during the first 24 hours. At least for the following 6 weeks, compression garments were used and strenuous activity was avoided.

Outcomes

The outcomes measured and analysed in this study included time to drain removal, total and daily volume of drain output, duration of hospital stay, emergency department visit, readmission to the hospital, secondary surgical procedure, and incidence of postoperative complications. The complications were defined as local or systemic. Systemic complications were defined as thromboembolic events, namely deep vein thrombosis or pulmonary thromboembolism. Local complications were defined as seroma, hematoma/bleeding, wound infection, wound dehiscence, and cutaneous necrosis. Seroma and hematoma were defined as a subcutaneous abdominal wall fluid collection evident on physical examination after drain removal that was successfully aspirated at least once (non-hematic clear fluid or hematic fluid, respectively); physical examination was considered suggestive of fluid collection when there was one of the following signs: erythema of the skin or scar, a visible distension, or a palpable tumefaction on the operated area with wave sign. If this was the case, a percutaneous puncture with a 21-gauge needle and a 20ml syringe was performed, for diagnosis and eventually for aspiration. Patients with liquid collections were examined within a week.

Drain output volume was registered daily, at the same time of day, by a nurse and all the patients were observed by one of the two surgeons daily until discharge. Patients were observed by one of the two surgeons at 1, 2, 3 and 4 weeks, and 2 and 3 months after surgery.

Statistical Analysis

Data were analysed using IBM SPSS version 21 software package (SPSS Inc.; Chicago, IL, USA). Normally distributed continuous variables were tested for normal

distribution and homogeneity of variance (Shapiro-Wilk and Levene tests respectively) and are presented as mean \pm standard deviation. Non-parametric continuous and ordinal variable is described as mean \pm standard deviation (median). Nominal categorical variables are described as percentages. T-Student and Mann-Whitney U tests were used to analyse continuous variables. χ^2 test was applied to analyse categorical variables. The probability level of 0.05 was used for rejection of the null hypothesis.

RESULTS

A total of 73 abdominoplasties with Scarpa fascia preservation were performed. In 33 patients, 2 closed-suction drains were used (group A) and in 40 patients, 3 closed-suction drains (group B).

The patients' general characteristics are summarized in Table 1 and did not differ significantly between groups. In group A, 7 patients had comorbidities, namely arterial hypertension, hypothyroidism (2 patients), venous insufficiency (2 patients), asthma, chronic obstructive pulmonary disease, and Crohn's disease; 22 patients had previous abdominal surgeries (c-section: 12 patients of which 7, 3 and 2 patients had one, two or three procedures; tubal ligation: 7 patients; total hysterectomy: 2 patients; appendectomy: 2 patients; hernia repair: 3 patients; and laparoscopic cholecystectomy: 2 patients); and 1 patient had previous bariatric surgery, having performed a laparoscopic sleeve gastrectomy. In group B, there were 4 patients with comorbidities, namely arterial hypertension (2 patients), Diabetes Mellitus type 1, and hypothyroidism; 18 patients had previous abdominal surgery (c-section: 13 patients, of which 5, 7 and 1 had one, two or three procedures; tubal ligation: 2 patients; total hysterectomy: 2

patients; oophorectomy: 1 patient; and hernia repair: 1 patient); and 2 patients had previous bariatric surgery (a laparoscopic Roux-en-Y gastric bypass and a laparoscopic adjustable gastric banding). In both groups, all the smokers reduced their consumption to 3 or less cigarettes per day 6 weeks prior to surgery.

Outcomes are summarized in Table 2 and Table 3. Figure 2 represents the daily evolution of drain output. Both, total and daily drain outputs, were lower in group B than in group A, although without significant statistical difference ($p>0.05$). Time until drain removal and duration of hospital stay are identical for every patient, since there was no other reason to prolong the hospitalization, being the patients discharged at the time of drain removal. None of the analysed outcomes had statistically significant differences between groups, even though group B tended to have a lower incidence of complications, namely seroma, hematoma, infection, wound dehiscence and DVT/PE. Group B had a higher incidence of necrosis and reoperation, but still without significant statistical difference.

In group A, complications were located as follows: among the 5 patients who developed seroma, 3 were located in the medial hypogastrium, 1 in the epigastrium and 1 in the umbilical region; hematomas were suprapubic (1 patient) or adjacent to the left drain perforation (1 patient); 1 patient had an infection in the umbilical region; wound dehiscence occurred mainly in the umbilical suture (5 patients), with 1 case in the inferior horizontal suture; cutaneous necrosis occurred in the umbilicus (1 patient); and 1 patient had an episode of deep venous thrombosis in the left leg.

In group B, complications were located as follows: 2 patients developed seroma in the epigastrium, 1 patient in the hypogastrium and 1 patient in the suprapubic region; 1 patient had a hematoma of the flank; no infections were identified; of the 4 patients

who developed wound dehiscence, 3 were located in the inferior horizontal suture, and 1 in the umbilical suture; cutaneous necrosis occurred in the umbilicus (2 patients); no vascular complications were reported.

It is important to highlight that all the fluid collections were of low volume (some as low as 15 ml of aspirated fluid) and resolved uneventfully with percutaneous aspirations in the office except for one case. This refers to a patient in group B who was reoperated during hospitalization, due to a hematoma of the flank, detected postoperatively, before hospital discharge. Neither of the groups registered emergency department visits or readmissions to the hospital. There were no deaths.

DISCUSSION

This prospective comparative study provides evidence that using 2 or 3 drains after abdominoplasty with Scarpa fascia preservation has no clinically important or statistically significant effects on the main outcomes (total and daily drain output, time to drain removal) or complications (local or systemic).

Closed-suction drains are still viewed as an important part of reconstructive and cosmetic surgery whenever dead space is created^{16, 17} and have been one of the most accepted and used means of prevention of wound complications in abdominoplasty.¹³ Being seroma the most common complication of abdominoplasty, the authors consider the use of drains, along with other precautions taken simultaneously, an advantage to reduce complications. As we have already pointed out, there are not many studies, either clinical or *in vitro*, analysing factors that may be relevant to improve suction drain efficiency. A systematic review on seroma prevention when potential spaces are

surgically created, as it is the case after a full abdominoplasty, has demonstrated that drains reduce seroma formation.¹⁶ The criteria used for drain removal is important for maximizing its efficiency. The utilization of a volume-dependent criteria for drain removal rather than a time-dependent criteria is more effective for seroma prevention¹⁶ considering a volume from 30 to 50 ml per 24 hours. Concerning intensity of drain vacuum, there are no studies published in the abdomen, but three randomized controlled trials involving 304 patients submitted to breast surgery, using two vacuum levels, high and low, did not show a significant effect of vacuum level in the analysed outcomes.²⁷⁻²⁹ Similar results were found in the present study, as group B operative field was exposed to a 50% higher total vacuum level than that of group A and no significant differences on the main outcomes or complications were observed. However, high-vacuum drainage (i.e. increased pressure differentials) has been proven to optimize fluid flow rate in an *in vitro* study.¹⁷ Despite the fact that there is no statistically significant differences between groups regarding total or daily drain output and seroma and hematoma incidences, there is a tendency for lower volumes and the fluid collections considered together (hematoma plus seroma) are 50% lower when three drains are used. A larger study population may clarify this issue.

One issue that is not addressed in most papers is the amount of fluid collected in the reservoir and the possible effect on drain efficiency. Other important issues which need to be clarified in clinical settings are: drain size and tubing length. A recent *in vitro* study analysed these variables and concluded that drain performance increases with perforated drains, increased intracavitary tubing length, decreased extracavitary tubing length and increased tubing diameter, being proportional to the negative pressure generated by the evacuator.¹⁷ The evacuator fill will have 50% less negative pressure as

it reaches 25% of its capacity.¹⁷ This may differ according to the evacuator volume.¹⁷ Nevertheless, it is something that should be taken into account by the surgeon using suction drains after an abdominoplasty. Clinical studies with patients submitted to a full abdominoplasty are needed to further clarify these important issues, preferably prospective, comparative and controlled.

This study was done with patients submitted to a full abdominoplasty with Scarpa fascia preservation, using bovie dissection with the same settings, and a criterion for drain removal of 50 ml per 24 hours per drain. When compared to a previously published randomized controlled trial by the same surgeons,²² who used dissection with an avulsion technique described by Vasconez³⁰ and a drain removal criteria of 30 ml per 24 hours per drain, we can verify an important evolution and advantage, since lower drain volumes (total an daily) and consequentially earlier drain removal were achieved. Indeed, time to drain removal was reduced from 3 to 2 days, representing an important enhancement of patient recovery and comfort. Nevertheless, the seroma rate increased from 2.5%, in the aforementioned study,²² to 10 to 15 % in the present study, for group A and B, respectively. This may be a consequence of the fact that drains were removed with a volume criterion of 50 ml per 24 hours. On the other hand, these were low volume seromas with no impact on the final result, in agreement with previous findings stating that seromas below 80 ml are not clinically problematic.¹³ Further studies, ongoing at our department, evaluating the effects of the dissection method and volumetric criteria for drain removal (30 *versus* 50 ml) on seroma rate incidence will allow for a better interpretation of these results.

This is the first prospective comparative clinical study on the number of drains in abdominoplasty with Scarpa fascia preservation. The clinical profile of the patients

included in this study is representative of the usual candidate for a full abdominoplasty - female, forty, with previous abdomen surgery, mainly c-section. Moreover, all surgeries were approached with the same surgical technique, using the same electrocoagulation device with the same settings, and with no concomitant procedures in the same operative time, by both surgeons working together during all surgeries, only differing on who was the main surgeon, performing a surgical technique and postoperative treatment protocol identical in all aspects, except the number of drains used.

CONCLUSIONS

Based on a comparative prospective study, our results suggest that using three closed-suction drains postabdominoplasty with Scarpa fascia preservation has no advantages in total and daily drain output, time to drain removal or complications (local or systemic) when compared with the usual two drains approach. Further clinical studies are essential to clarify other determinants of suction drain efficiency.

ACKNOWLEDGMENTS

The authors thank Marco Rebelo, M.D., for the help in collecting patient' data and collaborating with this study, and Isabel Bartosch, M.D., for providing the original drawing used in this article.

REFERENCES

1. Antonetti JW, Antonetti AR. Reducing seroma in outpatient abdominoplasty: analysis of 516 consecutive cases. *Aesthet Surg J* 2010; 30: 418-425.
2. Di Martino M, Nahas FX, Barbosa MV, et al. Seroma in lipoabdominoplasty and abdominoplasty: a comparative study using ultrasound. *Plast Reconstr Surg* 2010; 126: 1742-1751.
3. Pollock TA, Pollock H. Progressive tension sutures in abdominoplasty: a review of 597 consecutive cases. *Aesthet Surg J* 2012; 32: 729-742.
4. Neaman KC, Armstrong SD, Baca ME, et al. Outcomes of traditional cosmetic abdominoplasty in a community setting: a retrospective analysis of 1008 patients. *Plast Reconstr Surg* 2013; 131: 403e-410e.
5. Quaba AA, Conlin S, Quaba O. The no-drain, no-quilt abdominoplasty: a single-surgeon series of 271 patients. *Plast Reconstr Surg* 2015; 135: 751-760.
6. Costa-Ferreira A, Marco R, Vasconez L, et al. Abdominoplasty With Scarpa Fascia Preservation. *Ann Plast Surg* 2016; 76 Suppl 4: S264-274.
7. Grazer FM, Goldwyn RM. Abdominoplasty assessed by survey, with emphasis on complications. *Plast Reconstr Surg* 1977; 59: 513-517.
8. van Uchelen JH, Werker PM, Kon M. Complications of abdominoplasty in 86 patients. *Plast Reconstr Surg* 2001; 107: 1869-1873.
9. Kryger ZB, Fine NA, Mustoe TA. The outcome of abdominoplasty performed under conscious sedation: six-year experience in 153 consecutive cases. *Plast Reconstr Surg* 2004; 113: 1807-1817; discussion 1818-1809.

10. Kim J, Stevenson TR. Abdominoplasty, liposuction of the flanks, and obesity: analyzing risk factors for seroma formation. *Plast Reconstr Surg* 2006; 117: 773-779; discussion 780-771.
11. Stewart KJ, Stewart DA, Coghlan B, et al. Complications of 278 consecutive abdominoplasties. *J Plast Reconstr Aesthet Surg* 2006; 59: 1152-1155.
12. Neaman KC, Hansen JE. Analysis of complications from abdominoplasty: a review of 206 cases at a university hospital. *Ann Plast Surg* 2007; 58: 292-298.
13. Andrades P, Prado A, Danilla S, et al. Progressive tension sutures in the prevention of postabdominoplasty seroma: a prospective, randomized, double-blind clinical trial. *Plast Reconstr Surg* 2007; 120: 935-946; discussion 947-951.
14. Khan UD. Risk of seroma with simultaneous liposuction and abdominoplasty and the role of progressive tension sutures. *Aesthetic Plast Surg* 2008; 32: 93-99; discussion 100.
15. Matarasso A, Swift RW, Rankin M. Abdominoplasty and abdominal contour surgery: a national plastic surgery survey. *Plast Reconstr Surg* 2006; 117: 1797-1808.
16. Janis JE, Khansa L, Khansa I. Strategies for Postoperative Seroma Prevention: A Systematic Review. *Plast Reconstr Surg* 2016; 138: 240-252.
17. Khansa I, Khansa L, Meyerson J, et al. Optimal Use of Surgical Drains: Evidence-Based Strategies. *Plast Reconstr Surg* 2018; 141: 1542-1549.
18. Le Louarn C. [Partial subfascial abdominoplasty. Our technique apropos of 36 cases]. *Ann Chir Plast Esthet* 1992; 37: 547-552.
19. Le Louarn C. Partial subfascial abdominoplasty. *Aesthetic Plast Surg* 1996; 20: 123-127.

20. Costa-Ferreira A, Rebelo M, Vasconez LO, et al. Scarpa fascia preservation during abdominoplasty: a prospective study. *Plast Reconstr Surg* 2010; 125: 1232-1239.
21. Koller M, Hintringer T. Scarpa fascia or rectus fascia in abdominoplasty flap elevation: a prospective clinical trial. *Aesthetic Plast Surg* 2012; 36: 241-243.
22. Costa-Ferreira A, Rebelo M, Silva A, et al. Scarpa fascia preservation during abdominoplasty: randomized clinical study of efficacy and safety. *Plast Reconstr Surg* 2013; 131: 644-651.
23. Epstein S, Epstein MA, Gutowski KA. Lipoabdominoplasty without drains or progressive tension sutures: an analysis of 100 consecutive patients. *Aesthet Surg J* 2015; 35: 434-440.
24. Khan S, Teotia SS, Mullis WF, et al. Do progressive tension sutures really decrease complications in abdominoplasty? *Ann Plast Surg* 2006; 56: 14-20; discussion 20-11.
25. Bozola AR, Psillakis JM. Abdominoplasty: a new concept and classification for treatment. *Plast Reconstr Surg* 1988; 82: 983-993.
26. Matarasso A. Abdominolipoplasty: a system of classification and treatment for combined abdominoplasty and suction-assisted lipectomy. *Aesthetic Plast Surg* 1991; 15: 111-121.
27. van Heurn LW, Brink PR. Prospective randomized trial of high versus low vacuum drainage after axillary lymphadenectomy. *The British journal of surgery* 1995; 82: 931-932.
28. Bonnema J, van Geel AN, Ligtenstein DA, et al. A prospective randomized trial of high versus low vacuum drainage after axillary dissection for breast cancer. *Am J Surg* 1997; 173: 76-79.

29. Chintamani, Singhal V, Singh J, et al. Half versus full vacuum suction drainage after modified radical mastectomy for breast cancer- a prospective randomized clinical trial[ISRCTN24484328]. *BMC cancer* 2005; 5: 11.
30. Gardner PM, Vasconez LO. Liposculpture and lipectomy superficial to Scarpa's fascia. *Operative Techniques in Plastic and Reconstructive Surgery* 1996; 3: 42-46.

TABLES AND FIGURES

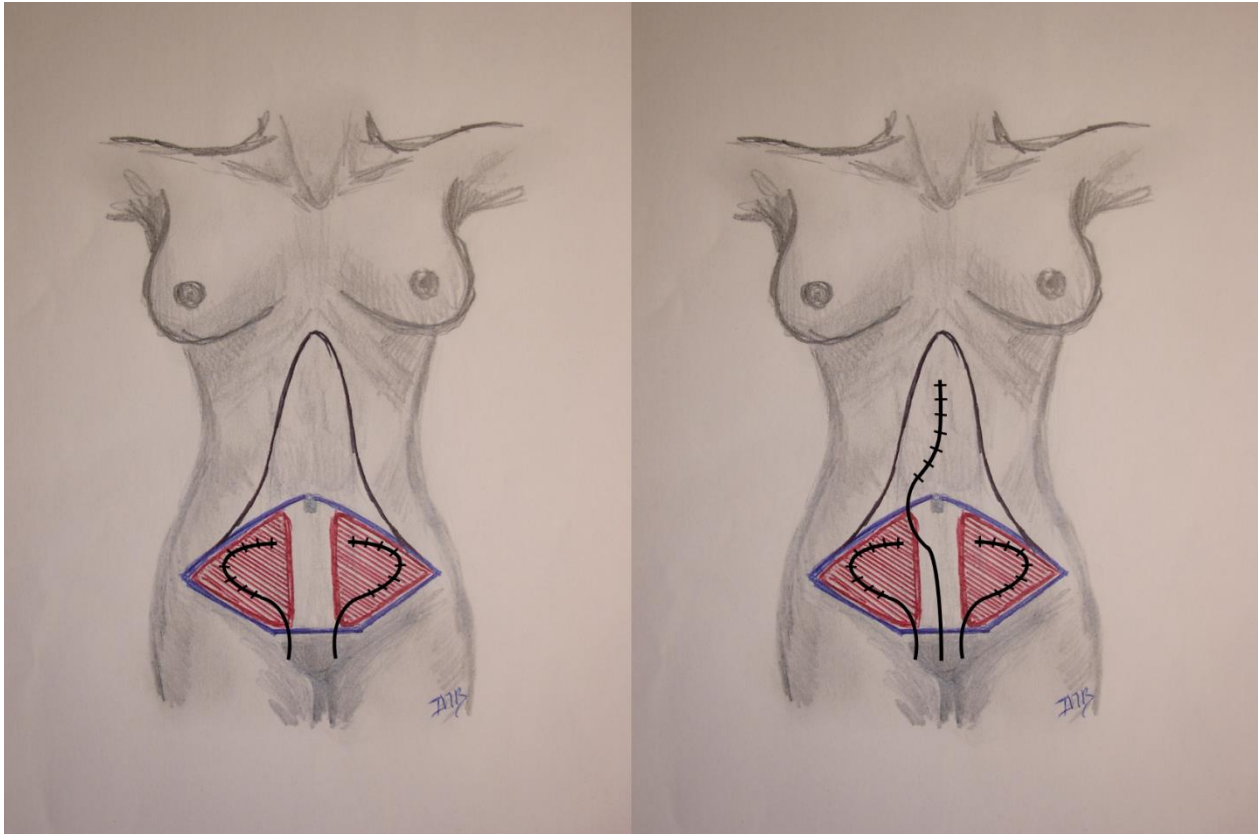


Figure 1. Representation of the closed-suction drains placement in group A (*left*) and group B (*right*). The *blue line* represents the skin resection. The *black line* limits the area to be undermined. The *red areas* show the region where dissection is performed on the plane of Scarpa fascia. The *curved lines* indicate the location of the closed-suction drains, with the traced portion representing the perforated part. (*Left*) In group A, two drains are placed, each in one iliac fossae, exiting in the lower hypogastrium, below the horizontal suture. (*Right*) In group B, a third drain is placed in the epigastric area, exiting alongside the other two.

Table 1. General characteristics of both Groups (n=73) *

	Group A	Group B	p - value
	(n=33)	(n=40)	
Age, years			
Mean ± SD	40.97± 8.76	41.20 ± 8.10	0.908
Range	25 – 61	22 – 57	
Body mass index, kg/m²			
Mean ± SD	25.40 ± 2.24	25.49 ± 3.32	0.896
Range	21.1 – 31.6	19.9 – 32.9	
Smoker			
Total number (%)	4 (12.1%)	5 (12.5%)	0.623
Comorbidities			
Total number (%)	7 (21.2%)	4 (10.0%)	0.158
Previous abdominal surgery			
Total number (%)	22 (66.7%)	18 (45.0%)	0.053
Previous bariatric surgery			
Total number (%)	1 (3.0%)	2 (5.0%)	0.573
Weight specimen, g			
Mean ± SD	853.03± 343.10	839.54 ± 420.03	0.741
Range	270.0 – 1630.0	140.0 – 1730.0	

*Group A: 2 drains (n=33), Group B: 3 drains (n=40).

Values presented as means ± standard deviation (SD) for data normally distributed and as percentages for nominal categorical variables. All the variables, except age, body mass index, and weight specimen, were compared between groups using the χ^2 test (Not significant; $p>0.05$). The other variables were compared using the Student's t-test (Not significant; $p>0.05$).

Table 2. Drain output of both Groups (n=73) *

	Group A (n=33)	Group B (n=40)	p - value
Time until drain removal, day			
Mean ± SD	2.30 ± 0.68 (2.0)	2.12 ± 0.40 (2.0)	0.138
Range	2.0 – 5.0	2.0 – 4.0	
Total drain output, ml			
Mean ± SD	204.09 ± 114.23	169.50 ± 101.07	0.315
Range	50.0 – 560.0	40.0 – 660.0	
Drain output day 1, ml			
Mean ± SD	129.39 ± 67.52	121.38 ± 74.34	0.634
Range	25.0 – 290.0	0.0 – 410.0	
Drain output day 2, ml			
Mean ± SD	54.70 ± 37.54	44.12 ± 30.99	0.192
Range	0.0 – 135.0	0.0 – 150.0	
Drain output day 3, ml			
Mean ± SD	14.39 ± 36.74	3.25 ± 12.28	0.076
Range	0.0 – 160.0	0.0 – 70.0	
Drain output day 4, ml			
Mean ± SD	3.64 ± 14.54	0.75 ± 4.74	0.241
Range	0.0 – 60.0	0.0 – 30.0	
Drain output day 5, ml			
Mean ± SD	1.82 ± 10.44	0.0 ± 0.0	0.274
Range	0.0 – 60.0	0.0 – 0.0	

*Group A: 2 drains (n=33), Group B: 3 drains (n=40).

Values presented as means ± standard deviation (SD) for data normally distributed and means ± standard deviation (median) for data not normally distributed (Time until drain removal). All the variables were compared between groups using the χ^2 test. Time until drain removal was analysed by the non-parametric test, Mann-Whitney U (Not significant; $p>0.05$).

Table 3. Outcomes of both Groups (n=73) *

	Group A (n=33)	Group B (n=40)	p - value
Seroma			
Total number (%)	5 (15.1%)	4 (10.0%)	0.377
Hematoma/bleeding			
Total number (%)	2 (6.1%)	1 (2.5%)	0.427
Infection			
Total number (%)	1 (3.0%)	0 (0.0%)	0.452
Wound dehiscence			
Total number (%)	6 (18.2%)	4 (10.0%)	0.251
Necrosis			
Total number (%)	1 (3.0%)	2 (5.0%)	0.573
DVT/PE			
Total number (%)	1 (3.0%)	0 (0.0%)	0.452
Emergency department visit			
Total number (%)	0 (0.0%)	0 (0.0%)	----
Readmission			
Total number (%)	0 (0.0%)	0 (0.0%)	----
Reoperation			
Total number (%)	0 (0.0%)	1 (2.5%)	0.548

DVT, deep venous thrombosis; PE, pulmonary embolism. Wound dehiscence refers to healing problems/suture rupture, without necrosis.

*Group A: 2 drains (n=33), Group B: 3 drains (n=40).

Values presented as percentages. All the variables were compared between groups using the χ^2 test (Not significant; $p>0.05$).

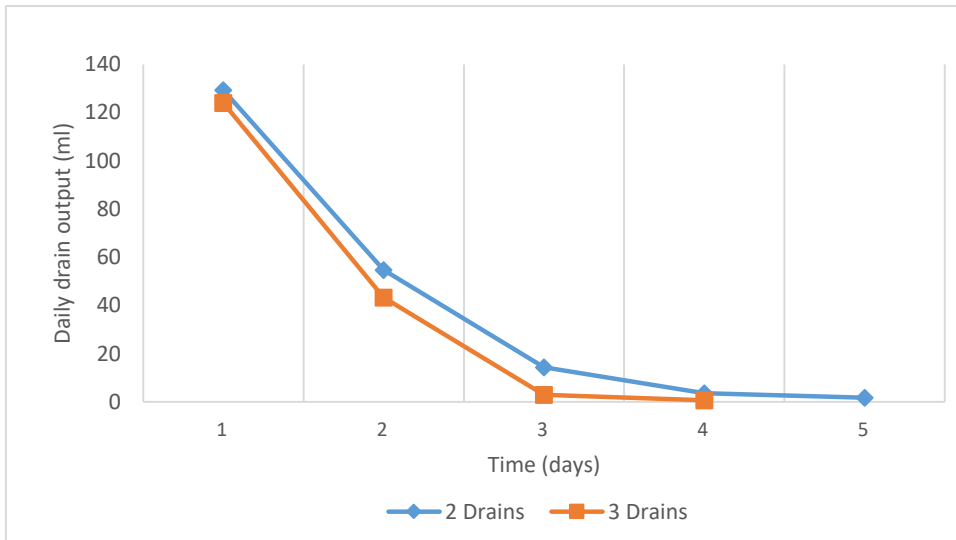


Figure 2. Average daily drain output (mean \pm SD) from group A (2 drains, n = 33) and group B (3 drains, n = 40) (Not significant; $p > 0.05$).

ANEXOS

Annals of Plastic Surgery

Online Submission and Review System

Author Resources

Instructions for Authors (this page)

[Reprint Ordering](#)

[Permissions Requests](#)

[Reprints](#)

Ethical/Legal Considerations

A submitted manuscript must be an original contribution not previously published (except as an abstract or preliminary report); must not be under consideration for publication elsewhere; and must, if accepted, not be published elsewhere in similar form, in any language, without the consent of Wolters Kluwer. Each person listed as an author is expected to have participated in the study to a significant extent. Although the editors and referees make every effort to ensure the validity of published manuscripts, the final responsibility rests with the authors, not with the Journal, its editors, or the publisher. The editorial office will acknowledge receipt of your manuscript and will give you a manuscript number for reference. Address all inquiries regarding manuscripts not yet accepted or published to the Journal's editorial office. **All manuscripts must be submitted online through the journal's website at <http://sap.edmgr.com>.** See submission instructions under "Online manuscript submission."

Patient anonymity and informed consent

It is the author's responsibility to ensure that a patient's anonymity be carefully protected and to verify that any experimental investigation with human subjects reported in the manuscript was performed with informed consent and following all the guidelines for experimental investigation with human subjects required by the institution(s) with which all the authors are affiliated. The protocol of the study must be approved by the Institutional Review Board (IRB) or the equivalent (eg, Research Ethics Board) where the study is conducted. Written releases from patients must accompany photographs in which the identity of the patient can be recognized. In the absence of such a release, an image must be cropped or partially obscured to the extent that the patient cannot be identified. Covering the eyes in a full-face photograph is not sufficient.

Conflicts of Interest and Copyright Transfer

Authors must state all possible conflicts of interest in the manuscript, including financial, consultative, institutional, and other relationships that might lead to bias or conflict of interest. If there is no conflict of interest, this should also be explicitly stated as "none declared." All sources of funding should be acknowledged in the manuscript. All relevant conflicts of interest and sources of funding should be included on the title page of the manuscript under the heading, "Conflicts of Interest and Source of Funding." For example:

Conflicts of Interest and Source of Funding: A has received honoraria from Company Z. B is currently receiving a grant (#12345) from Organization Y, and is on the speakers' bureau for Organization X – the CME organizers for Company A. For the remaining authors none were declared.

In addition, each author must complete and submit the journal's copyright transfer agreement, which includes a section on the disclosure of potential conflicts of interest based on the recommendations of the International Committee of Medical Journal Editors, "Uniform Requirements for Manuscripts Submitted to Biomedical Journals" (www.icmje.org/update.html).

A copy of the form is made available to the submitting author within the Editorial Manager submission process. Co-authors will automatically receive an e-mail with instructions on completing the form upon submission.

Compliance with NIH and Other Research Funding Agency Accessibility Requirements

A number of research funding agencies now require or request authors to submit the post-print (the article after peer review and acceptance but not the final published article) to a repository

that is accessible online by all without charge. As a service to our authors, Wolters Kluwer will identify to the National Library of Medicine (NLM) articles that require deposit and will transmit the post-print of an article based on research funded in whole or in part by the National Institutes of Health, Wellcome Trust, Howard Hughes Medical Institute, or other funding agencies to PubMed Central. The Copyright Transfer Agreement provides the mechanism.

Permissions

Authors must submit written permission from the copyright owner (usually the publisher) to use direct quotations, tables, or illustrations that have appeared in copyrighted form elsewhere, along with complete details about the source. Any permissions fees that might be required by the copyright owner are the responsibility of the authors requesting use of the borrowed material, not the responsibility of Wolters Kluwer.

Clinical Trials

Manuscripts based on a prospective clinical trial should document the registration of the clinical trial. If the trial is not registered, the authors should provide reasons for this omission. The final decision of the relevance of trial registration to any manuscript will be made by the editor.

Manuscript Submission

Online manuscript submission

All manuscripts must be submitted on line through the Web site: <http://sap.edmgr.com>.

First-time users: Please click the Register button from the menu and enter the requested information. On successful registration, you will be sent an e-mail indicating your user name and password. Print a copy of this information for future reference. Note: If you have received an e-mail from us with an assigned user ID and password, or if you are a repeat user, do not register again; simply log in. Once you have an assigned ID and password, you do not have to re-register, even if your status changes (that is, author, reviewer, or editor). **Authors:** Please click the log-in button from the menu at the top of the page and log into the system as an Author. Submit your manuscript according to the author instructions. You will be able to track the progress of your manuscript through the system. If you experience any problems, please contact: Jane Wood, Managing Editor, *Annals of Plastic Surgery*, e-mail: willhermyone@aol.com, or William C. Lineaweaver, MD, FACS, Editor-in-Chief, *Annals of Plastic Surgery*, e-mail: william.lineaweaver@jmsburncenters.com

Preparation of Manuscript

Manuscripts that do not adhere to the following instructions will be returned to the corresponding author for technical revision before undergoing peer review.

Articles submitted to the journal must be written with a solid basis of English language. If you need assistance in this area, listed below are a few companies who provide language and copyediting services. Use of an editorial service is at the discretion and cost of the authors, and will not guarantee acceptance for publication in the journal.

Please note: Appearance in the list of vendors does not represent endorsement by the publisher. Authors are encouraged to investigate each service on their own, as well as seek out additional vendors offering similar services.

- [American Journal Experts](#) (Discount available for Wolters Kluwer journal authors)
- [The Medical Editor](#)
- [Text Check](#)
- [Bio Science Writers](#)
- [Boston BioEdit](#)
- [ScienceDocs](#)

In addition, *Annals of Plastic Surgery* adheres to the SAMPL guidelines for statistical information. A link to the SAMPL guidelines may be found at <http://www.equator->

Each of the following should be submitted as a separate document within the submission file:

Cover letter: The cover letter should state the authors' intent to submit the article to *Annals of Plastic Surgery*, state the title of the article and authors' names, and contain any other information pertinent to the submission of the manuscript.

Title page: Include on the title page (a) complete manuscript title and a brief title for use as a running head; (b) authors' full names, highest academic degrees, and affiliations (limit of two); (c) name and address for correspondence, including fax number, telephone number, and e-mail address; (d) address for reprints if different from that of corresponding author; and (e) sources of support that require acknowledgment.

The title page must also include disclosure of funding received for this work from any of the following organizations: National Institutes of Health (NIH); Wellcome Trust; Howard Hughes Medical Institute (HHMI); RCUK; and other(s).

Structured or unstructured abstract and key words: Limit the abstract to 350 words. It must be factual and comprehensive. Limit the use of abbreviations and acronyms, and avoid general statements (eg, "the significance of the results is discussed"). The abstract should summarize the problem presented, studies undertaken, results, and conclusions; it replaces a summary at the end of the article.

Text: Provide succinct internal headings to clarify the paper's organization. Cite all tables and illustrations in the text. Define abbreviations at first mention in text and in each table and figure. If a brand name is cited, supply the manufacturer's name and address (city and state/country). Acknowledge all forms of support, including pharmaceutical and industry support, in an Acknowledgments paragraph.

Abbreviations: For a list of standard abbreviations, consult the Council of Biology Editors Style Guide (available from the Council of Science Editors, 9650 Rockville Pike, Bethesda, MD 20814) or other standard sources. Write out the full term for each abbreviation at its first use unless it is a standard unit of measure.

References: The authors are responsible for the accuracy of the references. Key the references (double-spaced) at the end of the manuscript. Cite the references in text in the order of appearance. Cite unpublished data, such as papers submitted but not yet accepted for publication or personal communications, in parentheses in the text. If there are more than three authors, name only the first three authors and then use et al. Refer to the List of Journals Indexed in *Index Medicus* for abbreviations of journal names, or access the list at <http://www.nlm.nih.gov/tsd/serials/lji.html>. Sample references are given below:

Journal article

1. Lin S-D, Tsai C-C, Lai C-S, et al. Endoscope-assisted parotidectomy for benign parotid tumors. *Ann Plast Surg* 2000;45:269-273

Book chapter

2. Todd VR. Visual information analysis: frame of reference for visual perception. In: Kramer P, Hinojosa J., eds. *Frames of Reference for Pediatric Occupational Therapy*. Philadelphia: Lippincott Williams & Wilkins; 1999:205-256

Entire book

3. Kellman RM, Marentette LJ. *Atlas of Craniomaxillofacial Fixation*. Philadelphia: Lippincott Williams & Wilkins; 1999

Software

4. Epi Info [computer program]. Version 6. Atlanta: Centers for Disease Control and Prevention; 1994

Online journals

5. Friedman SA. Preeclampsia: a review of the role of prostaglandins. *Obstet Gynecol* [serial online]. January 1988;71:22-37. Available from: BRS Information Technologies, McLean, VA. Accessed December 15, 1990

Database

6. CANCERNET-PDQ [database online]. Bethesda, MD: National Cancer Institute; 1996. Updated March 29, 1996

World Wide Web

7. Gostin LO. Drug use and HIV/AIDS [*JAMA HIV/AIDS* web site]. June 1, 1996. Available at: <http://www.ama-assn.org/special/hiv/ethics>. Accessed June 26, 1997

Supplemental Digital Content

Supplemental Digital Content (SDC): Authors may submit SDC via Editorial Manager to Wolters Kluwer journals that enhance their article's text to be considered for online posting. SDC may include standard media such as text documents, graphs, audio, video, etc. On the Attach Files page of the submission process, please select Supplemental Audio, Video, or Data for your uploaded file as the Submission Item. If an article with SDC is accepted, our production staff will create a URL with the SDC file. The URL will be placed in the call-out within the article. SDC files are not copy-edited by Wolters Kluwer staff; they will be presented digitally as submitted. For a list of all available file types and detailed instructions, please visit <http://links.lww.com/A142>.

SDC Call-outs: Supplemental Digital Content must be cited consecutively in the text of the submitted manuscript. Citations should include the type of material submitted (Audio, Figure, Table, etc.), be clearly labeled as "Supplemental Digital Content," include the sequential list number, and provide a description of the supplemental content. All descriptive text should be included in the call-out as it will not appear elsewhere in the article.

Example:

We performed many tests on the degrees of flexibility in the elbow (see Video, Supplemental Digital Content 1, which demonstrates elbow flexibility) and found our results inconclusive.

List of Supplemental Digital Content: A listing of Supplemental Digital Content must be submitted at the end of the manuscript file. Include the SDC number and file type of the Supplemental Digital Content. This text will be removed by our production staff and not be published.

Example:

Supplemental Digital Content 1. wmv

SDC File Requirements: All acceptable file types are permissible up to 10 MBs. For audio or video files greater than 10 MBs, authors should first query the journal office for approval. For a list of all available file types and detailed instructions, please visit <http://links.lww.com/A142>.

Digital figures:

A) Creating Digital Artwork

1. Learn about the publication requirements for Digital Artwork: <http://links.lww.com/ES/A42>
2. Create, Scan and Save your artwork and compare your final figure to the Digital Artwork Guideline Checklist (below).
3. Upload each figure to Editorial Manager in conjunction with your manuscript text and tables.

B) Digital Artwork Guideline Checklist

Here are the basics to have in place before submitting your digital artwork:

- Artwork should be saved as TIFF, EPS, or MS Office (DOC, PPT, XLS) files. High resolution PDF files are also acceptable.
- Crop out any white or black space surrounding the image.
- Diagrams, drawings, graphs, and other line art must be vector or saved at a resolution of at least 1200 dpi. If created in an MS Office program, send the native (DOC, PPT, XLS) file.
- Photographs, radiographs and other halftone images must be saved at a resolution of at least 300 dpi.
- Photographs and radiographs with text must be saved as postscript or at a resolution of at least 600 dpi.
- Each figure must be saved and submitted as a separate file. Figures should not be embedded in the manuscript text file.

Remember:

- Cite figures consecutively in your manuscript.
- Number figures in the figure legend in the order in which they are discussed.
- Upload figures consecutively to the Editorial Manager web site and enter figure numbers consecutively in the Description field when uploading the files.

Figure legends: Legends must be submitted for all figures. They should be brief and specific, and they should appear on a separate manuscript page after the references. Use scale markers in the image for electron micrographs, and indicate the type of stain used.

Color figures: The journal accepts for publication color figures that will enhance an article. Authors who submit color figures will receive an estimate of the cost for color reproduction. If they decide not to pay for color reproduction, they can request that the figures be converted to black and white at no charge.

Tables: Cite tables consecutively in the text, and number them in that order. Key each on a separate sheet, and include the table title, appropriate column heads, and explanatory legends (including definitions of any abbreviations used). Do not embed tables within the body of the manuscript. They should be self-explanatory and should supplement, rather than duplicate, the material in the text.

Style: Pattern manuscript style after the *American Medical Association Manual of Style* (10th edition). *Stedman's Medical Dictionary* (27th edition) and *Merriam Webster's Collegiate Dictionary* (11th edition) should be used as standard references. Refer to drugs and therapeutic agents by their accepted generic or chemical names, and do not abbreviate them. Use code numbers only when a generic name is not yet available. In that case, supply the chemical name and a figure giving the chemical structure of the drug. Capitalize the trade names of drugs and place them in parentheses after the generic names. To comply with trademark law, include the name and location (city and state in USA; city and country outside USA) of the manufacturer of any drug, supply, or equipment mentioned in the manuscript. Use the metric system to express units of measure and degrees Celsius to express temperatures, and use SI units rather than conventional units.

After Acceptance

Open access

Authors of accepted peer-reviewed articles have the choice to pay a fee to allow perpetual unrestricted online access to their published article to readers globally, immediately upon publication. Authors may take advantage of the open access option at the point of acceptance to ensure that this choice has no influence on the peer review and acceptance process. These articles are subject to the journal's standard peer-review process and will be accepted or rejected based on their own merit.

The article processing charge (APC) is charged on acceptance of the article and should be paid within 30 days by the author, funding agency or institution. Payment must be processed for the article to be published open access. For a list of journals and pricing please visit our [Wolters Kluwer Open Health Journals page](#).

Authors retain copyright

Authors retain their copyright for all articles they opt to publish open access. Authors grant Wolters Kluwer an exclusive license to publish the article and the article is made available under the terms of a Creative Commons user license. Please visit our [Open Access Publication Process page](#) for more information.

Creative Commons license

Open access articles are freely available to read, download and share from the time of publication under the terms of the [Creative Commons License Attribution-NonCommercial No Derivative \(CC BY-NC-ND\) license](#). This license does not permit reuse for any commercial purposes nor does it cover the reuse or modification of individual elements of the work (such as figures, tables, etc.) in the creation of derivative works without specific permission.

Compliance with funder mandated open access policies

An author whose work is funded by an organization that mandates the use of the [Creative Commons Attribution \(CC BY\) license](#) is able to meet that requirement through the available open access license for approved funders. Information about the approved funders can be found here: <http://www.wkopenhealth.com/inst-fund.php>

FAQ for open access

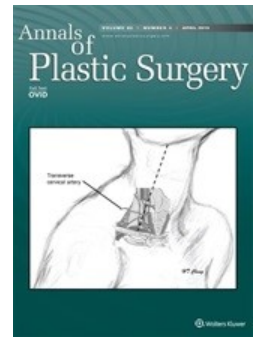
<http://www.wkopenhealth.com/openaccessfaq.php>

Page proofs and corrections: Corresponding authors will receive electronic page proofs to check the copyedited and typeset article before publication. Portable document format (PDF) files of the typeset pages and support documents (eg, reprint order form) will be sent to the corresponding author by e-mail. Complete instructions will be provided with the e-mail for downloading and printing the files and for faxing the corrected page proofs to the publisher. Those authors without an e-mail address will receive traditional page proofs. It is the author's responsibility to ensure that there are no errors in the proofs. Changes that have been made to conform to journal style will stand if they do not alter the authors' meaning. Only the most critical changes to the accuracy of the content will be made. Changes that are stylistic or are a reworking of previously accepted material will be disallowed. The publisher reserves the right to deny any changes that do not affect the accuracy of the content. Authors may be charged for alterations to the proofs beyond those required to correct errors or to answer queries. Proofs must be checked carefully and corrections faxed within 24 to 48 hours of receipt, as requested in the cover letter accompanying the page proofs.

Reprints: Authors will receive a reprint order form and a price list with the page proofs.

Reprint requests should be faxed with the corrected proofs, if possible. Reprints are normally shipped 6 to 8 weeks after publication of the issue in which the item appears. Contact the Reprint Department, Lippincott Williams & Wilkins/Wolters Kluwer, 351 W. Camden Street, Baltimore, MD 21201, fax: 410-528-4434, e-mail: reprints@wolterskluwer.com with any questions.

Publisher's contact: Fax corrected page proofs, reprint order form, and any other related materials to the Production Editor, at 877-705-1375.



Copyright © 2019 Wolters Kluwer Health, Inc. All rights reserved.

[Copyright/Disclaimer Notice](#) • [Privacy Policy](#)
