Comparison Between the Impact of Vasopressors and Goal-Directed Fluid Therapy on the Management of Free Flap Reconstruction of Head and Neck and Monitoring in ICU

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Introduction And Background

Free flap reconstruction of head and neck involves the microvascular heterotopic free tissue transfer including re-implantation of its blood supply [1]. This technique aims to restore function and cosmetic appearance of head and neck open wounds resulting from trauma or burns, and tumor excision with success survival rates exceeding 95% [2].

Increased risk of complications and higher cardiovascular morbidity and mortality frequently associate head and neck free flap reconstruction. Patients undergoing such procedures usually have low cardiovascular reserves that limit satisfactory organ perfusion with loss of function and even failure [3]. Therefore, ICU close monitoring for the first 24–48 hours post-operatively of the patient and fresh flap, anticoagulation, and fluid resuscitation is essential for the early prevention of complications [4].

Anticoagulants are used in preventing venous thromboembolism in high-risk patients immediately post-operatively especially when a patient’s Caprini risk score is > 8 while they should be stopped 24–48 hours post-operatively if the score is < 8. Caprini score is one of the further most valid in plastic head and neck reconstructive surgery [5]. Numerous anticoagulants such as aspirin, low molecular weight dextran, unfractionated heparin, prostaglandin-E1 are usually used. However, unfractionated heparin was associated with a high number of patients requiring revision surgery of the free flap’s anastomosis [6]. In order to optimize cardiac function and regional blood flow to the flap after adequate fluid delivery, vasopressors as epinephrine, norepinephrine, dobutamine, and dopexamine are usually useful [7].

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Recent review of vasopressor use showed a more than 80% lower rate of flap failure in patients [9]. Administration methods include both intravenous bolus and continuous infusion [8,9].

A continuously infused low-dose vasopressor drip in combination with the use of neuromuscular blocker to reduce the dose of systemic anesthesia can stabilize MAP and improve flap perfusion after anastomosis [10]. Bolus intravenous administration of vasopressors usually causes wide fluctuations in MAP producing unfavorable effect on the peripheral blood flow needed to perfuse the flap with long-lasting negative effects for its outcome. Therefore, clear coordination of the vasopressor timing, dosage, and agent must be implemented [11].

The intraoperative vasopressor of choice in maintaining normal blood pressure and flap blood flow in hypotensive patients is ephedrine, phenylephrine and norepinephrine [5-7]. Even though there is controversy regarding the use of intraoperative intravenous vasopressors as they induce vasoconstriction, thereby increasing the risk of postoperative thrombosis, decreased flap perfusion, and subsequent flap failure [12] with 70% of the surgeons prohibiting their intraoperative use [11-13]. Notably, these suppositions are chiefly grounded on experimental animal models with contradictory results [12,13]. Nevertheless, recent studies reported that intraoperative vasopressor use in free flap reconstruction did not alter their outcomes neither flap complications nor survival [7-10].

Notably, it was corroborated that there is no relationship between the timing of vasopressor administration either before, during, or after free flap reconstruction and postoperative failure rates [7-10]. On the other hand, no association was observed between the total dose of administered vasopressor and flap failure rates [9].

It has been hypothesized that local catecholamines are released due to sympathetic fibers activation during tissue dissection and free flap harvest [13]. Once the local catecholamines stream is washed-out, the acute hyperadrenergic stage is followed by a non-adrenergic stage with probable increase in collateral blood flow, and then by an increased adrenergic phase due to loss of moderating autonomic contribution in a delayed manner ranging from 48 hours to two weeks [14]. Therefore, the intraoperative use of vasopressors may actually increase flap perfusion due to enhanced overall mean arterial pressure without significant harm from sympathetomy [7,8].

There is an ongoing controversy on whether vasopressors have diverse effects on outcomes of bony versus soft tissue free flap reconstruction [11]. As corroborated in animal studies, bony free flaps typically involve longer procedures time causing increased flap ischemia and often depend on blood supply from periosteal perforators with a higher degree of ischemia relative to vasopressor administration, in contrast to soft tissue free flaps [15].

Review

Physiological effects of vasopressors

The homeostasis of the body is mainly controlled by the autonomic nervous system, including parasympathetic and sympathetic components. The adrenergic α- and β-receptors are responsible for the vascular tone and cardiovascular function. The α-1 agonists help smooth muscle contraction, resulting in increased systemic vascular resistance and MAP [13]. On the contrary, α-2 agonists cause smooth muscle relaxation and contribute to platelet aggregation through the activation of α-2 receptors on platelets. The positive inotropic and chronotropic effects of β-1 agonists mainly increase cardiac output, while β-2 agonists cause smooth muscle relaxation in the lungs [16].

Ephedrine has lesser effects on peripheral vasoconstriction as it is an indirect sympathetic agonist with primarily strong β-1 and β-2 effects and weak α-1 stimulation. Hypotension is considered when the MAP became lower than 60 mmHg as the arterial blood pressure during anastomosis should be equal to or more than 70 mmHg.

The significant principle of optimizing cardiac performance to enhance free flap perfusion is the goal of fluid management. Based on the current evidence, perioperative goal-directed therapy can improve postoperative outcome of intermediate-to-high-risk surgical patients. It seems to be associated with decreased postoperative length of stay, complications and possibly even mortality. Fluid resuscitation has been chosen for hemodynamic support after free flap ischemia from systemic vasopressors [17].

Significant amounts of intraoperative intravenous fluids are administrated to avoid intraoperative vasopressor use. Fluid resuscitation always carries the risk of both limited and excessive fluid delivery resulting in flap failure with end-organ dysfunction [18].

Colloids should be used cautiously in free flap patients. Patients should be kept hydrated for the sake of improved flap survival. Crystalloid fluid resuscitation is used to ensure adequate flap and renal perfusion with a goal urine output of 0.5-1.0 mL/kg/hour [19].
Crystallloid volumes should not exceed 130 mL/kg per 24-hour period. The controversy about colloid versus crystallloid solutions in fluid resuscitation has long been discussed in the recent years and ended with the favor of crystalloids. A large meta-analysis by Perel et al. accomplished that colloids are significantly more expensive than crystalloids and some of them might even increase mortality rather than reducing it [20].

Greater total intravenous fluid volume of crystalloids than colloids are needed to meet the same hemodynamic stability [19]. The intravenous fluids (IVF) administration is the first-line treatment that combats acute changes in blood pressure, whereas, their excessive administration may result in fluid overload and decreased flap microcirculation via hemodilution-induced hypercoagulability contributing to free flap failure [2,7,18].

Free flaps edema is mainly owed to flap denervation that leads to absent lymphatic drainage pathways and poor interstitial fluid reabsorption [7]. Furthermore, deleterious flap and patient outcomes may occur secondary to pulmonary edema and associated cardiopulmonary complications resulting from excessive fluid resuscitation [10-18].

On the other hand, increased free flap complications can also result from under-resuscitation due to decreased flap perfusion, deteriorated by further hypotension from anesthetic agents. Ischemia-reperfusion injury can aggravate inflammation with subsequent flap failure [21]. Therefore, the microvascular surgeons and anesthesiologists must be vigilant in the administration of vasopressors and IVF to improve free flap and patient outcomes. Of note, the use of vasopressors may be an advantageous alternative to IVF to regulate systemic perfusion pressure and flap perfusion intraoperative [5].

It is well known that head and neck free flap reconstruction requires long operative and hospitalization times [22]. Goal-directed fluid replacement (equal weight in the immediate pre- and postoperative period) is one of the protocols specific to head and neck surgery that was designed to optimize patients' cardiovascular performance and thus proposed to reduce surgical morbidity and reduce the length of stay by utilizing a multimodal and multidisciplinary algorithms [6–21].

Other enhanced recovery protocols include standard perioperative management, especially pre-operative assessment, thermoregulation, hemodynamic and flap monitoring, deep vein thrombosis (DVT) prophylaxis, analgesia and antibiotics guidelines, intensive care unit admission and early mobilization [6].

The goal-directed fluid replacement is mainly dependent on the tissue perfusion physiology. For example, it modulates the adequate oxygen supply to tissues either by its content in blood or the blood flow or depressing organ demands [23]. Although the local tissue blood flow is dependent on systemic blood pressure, it is autoregulated to keep constant blood supply under a wide range of blood pressure. The best example of this hypothesis in case of severe hypoperfusion is the redistribution of blood flow to vital organs (i.e., heart and brain) while gastrointestinal tract, kidneys, skin, etc. may suffer hypoperfusion (occult hypoperfusion). Consequently, controlling cardiac index remains the basis of goal-directed fluid replacement which can be achieved by functional hemodynamic monitoring [22]. Conferring to these data, surgical wound healing is associated with increased tissue oxygen consumption which necessitates an increase of the cardiac output and modulation of systemic vascular resistance in order to pass the perioperative period without organ failure [21].

Goal-directed fluid replacement is possible by modulating any of the parameters affecting stroke volume (SV)-preload, contractility or afterload. For instance, increasing the preload can be achieved using IVF replacement and increasing heart contractility can be attained by inotropes (dobutamine or dopexamine) but with the cost of increased myocardial oxygen consumption. Lastly, afterload is usually moderated to reach adequate perfusion pressures, but with local flow redistribution. Consequently, goal-directed fluid replacement is highly recommended in high-risk surgical patients [24].

Although fluid management has been directed by monitoring standard vital signs such as blood pressure, heart rate and urine output, these endpoints are insensitive to small changes in hemodynamic stability and hypoperfusion [14].

Recently, goal-directed therapy (GDT) is based on the Frank-Starling mechanism which states that the heart contractility increases with greater diastolic filling. When the total circulating blood volume is increased by intravenous fluids administration, cardiac output is thus increased. On the other hand, if hypovolemia occurs, this cardiac output augmentation will reach a plateau, then decreases [21].

In GDT, the intravenous fluids, vasopressors and inotropic administration are guided by instantaneous measurements of stroke volume (SV) and stroke volume variation (SVV) in one respiratory cycle to optimize cardiac output (CO). If hypovolemia occurs, the SVV will be exaggerated, with more response to fluid administration and better hemodynamic stability [22]. In order to perform GDT, the FloTrac/Vigileo system (Edwards Lifesciences LCC, Irvine, CA) is used for monitoring hemodynamic stability. Vigileo monitor continuously analyses the arterial pressure waveform to calculate SV, SVV, CO and cardiac index (CI; CO / total body surface area) through a connected transducer in a radial arterial line [25,26].
To the best of our knowledge, the SVV-guided hemodynamic treatment keeping SVV ≤ 13% improves oxygen delivery, reduces intravenous fluids perioperatively, as well as medical complications and hospital stay compared to conventional fluid management in head and neck patients [27,28].

Predictors of flap failure
Flap failures comprise pre-operative patient characteristics as well as peri- and post-operative factors. Pre-operative aspects including medical status (cardiovascular disease, stroke, asthma, diabetes, chronic obstructive pulmonary disease and renal disease), age, body mass index, smoking, previous blood transfusions, comorbid conditions, chemotherapy and radiation therapy are paramount determinants of outcome [29].

Perioperative variables were collected from intraoperative time, number of flaps, and type of reconstruction, CO monitor, volume and type of administered intravenous fluid, blood transfusion, blood loss, urine output, and administered vasoconstrictors or inotropes [22].

Postoperative variables are the duration of ICU hospitalization, 30-day flap survival and flap-related complications including flap dehiscence, fistula, donor site complications, need for reoperation due to flap failure [14-30].

The major features contributing to free flap failure involve mainly the condition of the tissue, mechanical compression, as well as fibrin dominant venous and platelet dominant arterial thrombosis. These facts indicate using antiplatelet agents or heparin to avoid arterial and venous thrombosis respectively. Surgical manipulations contribute mainly to tissue condition. Outside the native excellence of the flap, rough management of the tissues at the site of injury leads to deteriorating edema and inflammation with associated hypercoagulability [14].

Previous animal studies that examined vasopressor administration in free flaps have reported that flaps were more subtle to vasopressors once detached from the innate nervous structure of the original site of donor tissue aggravating flap failure. Mechanical compression varies according to patient movement, surgical tailoring, or hematoma following uncontrolled and recurrent bleeding [31].

Monitoring of head and neck free flaps
Moreover, it was proven that near-infrared spectroscopy (NIRS) can continuously measure free flap tissue oxygenation and perfusion hence predicting microvascular thrombosis in free flaps even before clinical signs, allowing great opportunity for flap rescue [32].

It is a non-invasive well-accepted modality that evaluates flap viability at the bedside directing attention to impending flap failure, allowing prompt surgical intervention. Another modality that evaluates oxygen saturation percent (StO2%) in the tissues beneath the sensor is tissue oximetry that uses two wavelengths of 690 and 830 nm near-infrared light which spread to the tissue, and then collected by four photo detectors located in each sensor. Measurements can be affected by the probe position [32].

NIRS is more advantageous than laser Doppler modality in identifying arterial flow complications as its values are not affected by the patient movement [33]. Also, NIRS has the advantage of light capability to penetrate deeply into large volume of tissues in a non-invasive, cheap, reliable and reproducible manner. However, laser Doppler flowmetry can only observe superficial microcirculatory changes in tissue [34].

Another promising additional parameter is the new Hypotension Probability Indicator that could be advantageous to discover any drop in the MAP, before hypotension occurs [35].

Conclusions
Recently, GDFT has been proven to be excellent in high-risk head and neck free tissue transfer surgery as it decreases prolonged intensive care unit admission (ICU) hospitalization and complication rate. Today, GDFT is highly suggested as one of the enhanced recovery after surgery protocols for major head and neck free flap reconstruction surgery.

Additional Information
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References

1. Hagans N, Longgro D: Anesthesia for free vascularized tissue transfer. Microsurgery. 2009, 29:161-167. 10.1002/micr.20584

2. Haughey BH, Wilson E, Klueve L, et al.: Free flap reconstruction of the head and neck: analysis of 241 cases. Otalaryngol Head Neck Surg. 2001, 125:10-17. 10.1067/mhn.2001.116788

3. Pereira CMJ, Figueiredo MEL, Carvalho R, Carte D, Assunção JP: Anesthesia and surgical microvascular flaps. Braz J Anesthesiol. 2012, 62:563-579. 10.1590/S0034-7094(2012)07156-3

4. Swartz JE, Aarts MC, Smart WM, et al.: The value of postoperative anticoagulants to improve flap survival in the free radial forearm flap: a systematic review and retrospective multicentre analysis. Clin Otolaryngol. 2015, 40:600-609. 10.1111/coa.12425

5. Wax MK, Azzi I: Perioperative considerations in free flap surgery: a review of pressors and anticoagulation. Oral Oncol. 2018, 83:154-157. 10.1016/j.oraloncology.2018.06.025

6. Coyle MJ, Main B, Hughes C, et al.: Enhanced recovery after surgery (ERAS) for head and neck oncology patients. Clin Otolaryngol. 2016, 41:118-126. 10.1111/coa.12482

7. Geh CSL, Ng MM, Song DH, Ooi ASH: Perioperative vasopressor use in free flap surgery: a systematic review and meta-analysis. J Reconstr Microsurg. 2019, 35:259-340. 10.1055/s-0039-1687914

8. Rose L, Prado N, Muvey D, Laugharne D, Jones K: Vasoactive agents do not adversely influence the success of maxillofacial microvascular free-flap surgery: a retrospective analysis. Int J Anesthesiol Res. 2016, 4:327-330. 10.19070/2352-2780-160067

9. Monroe MM, Canary SD, Ghunem TA, Swide CE, Wax MK: Safety of vasopressor use in head and neck microvascular reconstruction: a prospective observational study. Otolaryngol Head Neck Surg. 2011, 144:877-882. 10.1177/01945998114013113

10. Fang L, Liu J, Yu C, Hanasono MM, Zheng G, Yu P: Intraoperative use of vasopressors does not increase the risk of free flap compromise and failure in cancer patients. Ann Surg. 2018, 268:579-384. 10.1097/SLA.0000000000002295

11. Motakef D, Mountzardis PM, Ismail IK, Agag RL, Patel A: Perioperative management for microsurgical free tissue transfer: survey of current practices with a comparison to the literature. J Reconstr Microsurg. 2015, 31:355-363. 10.1055/s-0035-1546422

12. Ibrahim AMS, Kim PS, Rabie AN, Lee BT, Lin SJ: Vasopressors and reconstructive flap perfusion: a review of the literature comparing the effects of various pharmacologic agents. Ann Plast Surg. 2014, 75:245-248. 10.1097/SAP.0b013e31828d70b3

13. Eley KA, Young JD, Watt-Smith SR: Power spectral analysis of the effects of epinephrine, norepinephrine, dobutamine and dopexamine on microcirculation following free tissue transfer. Microsurgery. 2015, 35:275-281. 10.1002/micr.22072

14. Eskander A, Kang S, Twell B, et al.: Predictors of complications in patients receiving head and neck free flap reconstructive procedures. Otolaryngol Head Neck Surg. 2018, 158:859-847. 10.1016/j.earl.2018.09.083

15. Driessens M, Vanhoutte PM: Vascular reactivity of the isolated tibia of the dog. Am J Physiol. 1979, 236:H904-H908. 10.1152/ajpheart.1979.236.6.H904

16. Pardo MC, Miller R: Basics of anesthesia. Elsevier, Philadelphia, PA; 2018.

17. Vyas K, Wong L: Intraoperative management of free flaps: current practice. Ann Plast Surg. 2014, 72:S213-S219. 10.1097/SAP.0000000000000096

18. Ettinger KS, Arce K, Lohse CM, et al.: Higher perioperative fluid administration is associated with increased rates of complications following head and neck microvascular reconstruction with fibular free flaps. Microsurgery. 2017, 37:128-136. 10.1002/micr.30061

19. Cortés DO, Barros TG, Njimi H, Vincent JL: Crystalloids versus colloids: exploring differences in fluid requirements by systematic review and meta-regression. Anesth Analg. 2015, 120:369-402. 10.1213/ANE.0000000000001564

20. Perel P, Roberts I, Ker K: Colloids versus crystalloids for fluid resuscitation in critically ill patients. Cochrane Database Syst Rev. 2015, 2015:CD005567. 10.1002/14651858.CD005567.pub6

21. Dott JC, Farwell DG, Findlay M, et al.: Vasoactive agents do not adversely influence the success of microvascular reconstruction: a prospective observational study. J Reconstr Microsurg. 2015, 31:355-363. 10.1055/s-0039-1687914

22. Wetmore KM, Egan TD, Moneta G, et al.: Goal-directed intraoperative fluid administration reduces length of hospital stay after major surgery. Anesthesiology. 2002, 97:820-826. 10.1097/00000542-200201000-00012

23. Chalmers A, Turner MWH, Anand R, Puxeddu R, Brennan PA: Cardiac output monitoring to guide fluid replacement in head and neck microvascular free flap surgery-what is current practice in the UK? Br J Oral Maxillofac Surg. 2012, 50:500-505. 10.1016/j.bjoms.2011.08.010

24. Eldredge M, Esposito I, Kassim D: Outcome of intraoperative goal-directed therapy using Vigileo/FloTrac in high-risk patients scheduled for major abdominal surgeries: a prospective randomized trial. Egyptian J Anaesth. 2017, 33:263-269. 10.1016/j.eja.2017.05.002

25. Michaels F: Changes in arterial pressure during mechanical ventilation. Anesthesiology. 2005, 105:419-428. 10.1097/00000542-200508000-00026

26. Scheeren TWJ, Wiesnack C, Gerlach H, Marx G: Goal-directed intraoperative fluid therapy guided by stroke volume and its variation in high-risk surgical patients: a prospective randomized multicentre study. J Clin Monit Comput. 2013, 27:225-233. 10.1007/s10571-013-9461-6

27. Clark JR, McCluskey SA, Hall F, et al.: Predictors of morbidity following free flap reconstruction for cancer
30. Patel RA, McCluskey SA, Goldstein DP, et al.: Clinicopathologic and therapeutic risk factors for perioperative complications and prolonged hospital stay in free flap reconstruction of the head and neck. Head Neck. 2010, 32:1345-1353. 10.1002/hed.20639
31. Gardiner MD, Nanchahal J: Strategies to ensure success of microvascular free tissue transfer. J Plast Reconstr Aesthet Surg. 2010, 63:665-673. 10.1016/j.bjps.2010.06.011
32. Payette JR, Kohlenberg E, Leonardi L, et al.: Assessment of skin flaps using optically based methods for measuring blood flow and oxygenation. Plast Reconstr Surg. 2005, 115:539-546. 10.1097/01.PRS.0000148415.54546.CA
33. Hölzle F, Loeffelbein DJ, Nohte D, Wolff K-D: Free flap monitoring using simultaneous non-invasive laser Doppler flowmetry and tissue spectrophotometry. J Craniomaxillofac Surg. 2006, 34:25-33. 10.1016/j.jcms.2005.07.010
34. Noblett SE, Snowden CP, Shenton BK, Horgan AF: Randomized clinical trial assessing the effect of Doppler-optimized fluid management on outcome after elective colorectal resection. Br J Surg. 2006, 93:1069-1076. 10.1002/bjs.5454
35. Calvo-Vecino JM, Ripollés-Melchor J, Mythen MG, et al.: Effect of goal-directed haemodynamic therapy on postoperative complications in low-moderate risk surgical patients: a multicentre randomised controlled trial (FEDORA trial). Br J Anaesth. 2018, 120:734-744. 10.1016/j.bja.2017.12.018