

Identification of Independent Risk Factors for Complications: A Retrospective Analysis of 163 Fibular Free Flaps for Mandibulofacial Reconstruction

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Purpose: Fibular free flap transfer is a powerful tool available to the reconstructive surgeon when treating oral and maxillofacial defects, but complications still occasionally occur and predictive analysis focusing on this specific flap is limited in terms of risk factors for complication. The purpose of this study was to identify key variables associated with complications in patients undergoing fibular free flap transfer.

Patients and Methods: The data of 163 consecutive patients who underwent fibular free flap surgery at the Department of Oral and Maxillofacial Surgery, Sun Yat-Sen Memorial Hospital of Sun Yat-Sen University, between 2012 and 2015 were reviewed retrospectively. Patient demographic data, laboratory data, surgical data, and fluid infusion-related data that may have an influence on free flap outcomes were recorded. Univariate and multivariate logistic regression analyses were used to identify relevant risk factors.

Results: A total of 163 fibular free flaps were transferred for mandibulofacial reconstruction in 163 patients with a mean age of 50.9 years. Postoperative complications developed in 33 (20.2%). Multivariate analysis showed that free flap complications were significantly associated with radiotherapy history (odds ratio [OR], 5.12; $P = .001$), postoperative anemia (OR, 1.048; $P = .041$), postoperative hypoalbuminemia (OR, 0.844; $P = .002$), and prolonged operative time (OR, 1.005; $P = .004$).

Conclusions: Radiotherapy history, decreased postoperative hemoglobin and albumin levels, and prolonged operative time are potential predictors of postoperative complications after fibular free flap reconstruction for mandibulofacial defects.

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113 With advancements in reconstructive microsurgery,
 114 fibular free flap transfer has become the preferred
 115 reconstructive technique at many oncology centers
 116 Q3 to repair complex defects of the maxilla and mandible.
 117 Although various donor sites are available to provide
 118 vascularized bone grafts for reconstruction, including
 119 the fibular flap, radius flap, iliac flap, and scapular flap,
 120 the fibular free flap has been considered the flap of first
 121 choice for the reconstruction of mandibular continuity
 122 defects in patients with oral cavity cancer,¹ owing to
 123 its potential for contouring, the implant osseointegration,
 124 the satisfying pedicle length, and the skin paddle
 125 reliability.² However, as reported, reconstructions of
 126 head and neck sites were significantly associated
 127 with higher rates of flap failure compared with other
 128 Q4 sites such as the breast and extremities.³ Although surgical
 129 technical issues are dominant factors that
 130 contribute to microvascular free flap failure, many
 131 nontechnical variables are also contributory. In fact,
 132 predictors of complications after reconstruction of
 133 mandibular continuity defects with fibular free flaps
 134 have been extensively studied. However, most of the
 135 research analyzed different types of flaps at the same
 136 time, which compromises the specificity and applicability
 137 of their outcomes in fibular free flap microsurgery.^{4,5}
 138 Moreover, perioperative fluid management, which is an
 139 important aspect of patient care, has not been adequately
 140 explored in fibular flap transfer. There is currently a
 141 dearth of guidelines regarding the optimal rate or volume
 142 of perioperative fluid infusion, but it has been well
 143 shown that fluid therapy plays a role in microsurgical
 144 outcomes of head and neck free flap reconstruction,
 145 and aggressive fluid delivery has been found to be an
 146 independent predictor of complications after free
 147 flap reconstruction of the head and neck.⁶ However,
 148 to our knowledge, there has been no literature assessing
 149 fluid management solely in fibular free flap microsurgery.
 150 For these reasons, it may be more prudent to explore
 151 risk factors for complications in a study that includes
 152 only fibular free flaps.

153 Therefore, we undertook a retrospective study
 154 focusing on fibular free flap reconstruction for mandibulofacial
 155 defects. We hypothesized that some treatment-related
 156 variables such as the intraoperative infusion rate of fluid
 157 and operative time could influence the outcome. The
 158 specific aim of the study was to estimate the effect of
 159 demographic and treatment-related variables on the
 160 development of early in-hospital complications.

164 Patients and Methods

165 STUDY DESIGN

166 To address the research purpose, we designed and
 167 implemented a retrospective review of fibular free
 168

169 flaps in mandibulofacial reconstruction surgery. The
 170 study protocol was approved by the Institutional Review
 171 Board (IRB) of Sun-Yat San Memorial Hospital of Sun-Yat
 172 San University. Because the work was designed to
 173 retrospectively review medical records and was certified
 174 by the IRB as low risk, informed consent was not
 175 required by the IRB. The medical records Q5 of all
 176 consecutive patients who underwent fibular free flap
 177 interventions for reconstruction of defects in the
 178 mandibulofacial region at Sun Yat-Sen Memorial Hospital
 179 (Guangzhou, China) between January 1, 2012, and
 180 December 31, 2015, were reviewed.

182 POTENTIAL PREDICTOR VARIABLES

183 Potential predictors were selected based on review
 184 of the literature⁷ and clinical experience. Demographic,
 185 surgical, and fluid infusion data were recorded. Demographic
 186 variables included age, tobacco use, comorbidities,
 187 history of radiotherapy, reasons for flap, and preoperative
 188 and postoperative hemoglobin (Hgb) and albumin levels.
 189 Active smoking Q6 was determined at the time of diagnosis;
 190 hence, patients who ceased smoking after primary
 191 referral were counted as active. Regarding comorbidities,
 192 heart disease, hypertension, and diabetes were investigated
 193 because they could have altered the condition of the
 194 peripheral blood vessels.⁸ Surgical variables included
 195 intraoperative blood loss, intraoperative blood transfusions,
 196 and duration of surgery. A blood transfusion was indicated
 197 when the Hgb level was lower than 7 g/dL or the hematocrit
 198 (Hct) level was lower than 25% in patients with
 199 uncompromised function (cardiac or pulmonary). In
 200 hemodynamically Q7 stressed patients, a blood transfusion
 201 was recommended when the Hct level was lower than
 202 25% for patients younger than 60 years or when the
 203 Hct level was lower than 30% for patients older than
 204 60 years.⁹ Fluid variables included the intraoperative
 205 intravenous crystalloid and colloid infusion rate and
 206 the intravenous crystalloid and colloid infusion rate
 207 in the first 24 hours postoperatively, both of which were
 208 standardized to the patient's body weight (in milliliters
 209 per kilogram per hour). The intraoperative fluid
 210 infusion was determined at the discretion of the
 211 anesthesiologists on the basis of intra-arterial blood
 212 pressure monitoring, the stroke volume variation, and
 213 the patient's urine output. The rate of postoperative
 214 fluid infusion was titrated by the surgical team, taking
 215 into account the patient's heart rate, blood pressure,
 216 and urine output.

220 OUTCOME MEASURES

221 The primary outcome was postoperative in-hospital
 222 complications defined as any adverse event requiring
 223 intervention or affecting length of stay. We
 224

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