



# The role of robotic surgery in emergency setting

## Study Protocol

<b>Title:</b>	The role of robotic surgery in emergency setting
<b>Sponsor:</b>	World Society of Emergency Surgery (WSES)
<b>Coordinator</b>	Marco Milone, MD, PhD, FACS Associate Professor of Surgery
<b>Proposing Institution</b>	University of Naples “Federico II” Department of Clinical Medicine and Surgery
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<b>Rationale</b>	<p>Robotic surgery has progressively gained acceptance in several surgical fields, being routinely used for elective interventions<sup>1-3</sup>. The issue regarding the role of robotic surgery for emergency procedures remains open. Few studies have been published regarding the applications of robotics for emergency general surgery procedures; they were reviewed and discussed in the 2021 WSES position paper<sup>4</sup>. Studies on colorectal surgery, hiatal hernia surgery, bariatric surgery, gallbladder surgery, and abdominal wall surgery were included and statements proposed. The experts recommended a strict patient selection, an adequate training of the operating surgical team and an improvement of the accessibility of the robotic platforms. We propose this prospective study to better define the application of robotic surgery in an emergency setting, evaluating the intraoperative and postoperative outcomes, trying to understand the role of the robotic platform in the management of emergency situations.</p>

<b>Aims of the study</b>	Evaluate safe and feasibility of robotic surgery in emergency setting.
<b>Clinical Phase:</b>	Observational, prospective, multicentre
<b>List of participating Centres</b>	TBD
<b>Study design:</b>	<p><b>Data of clinically stable patients who underwent robotic surgery in emergency setting will be prospectively analysed.</b> The pathologies that will mainly be taken into consideration will be acute diverticulitis, acute cholecystitis and obstructed hernias. The Hinchey classification will be used to describe the degree of acute diverticulitis <sup>5</sup>, and the 2018 Tokyo guidelines will be used to describe the degree of acute cholecystitis <sup>6</sup>. Patients with other surgical pathologies may also be enrolled in the study as long as they are treated in robotic surgery in emergency setting.</p> <p><b>Data relating to the operating theatre team and the surgical instruments used will be collected in order to conduct a cost analysis.</b></p> <p>Data will be collected in a designated database.</p>
<b>Inclusion Criteria</b>	<ul style="list-style-type: none"> <li>• Age &gt; 18 years old</li> <li>• Clinically stable patients with disease requiring emergency surgical treatment</li> <li>• Intervention performed in robotic surgery</li> <li>• Capability of giving valid informed consent</li> </ul>
<b>Exclusion Criteria</b>	<ul style="list-style-type: none"> <li>• Age &lt; 18 years old</li> <li>• Intervention performed in open or laparoscopic surgery</li> <li>• Elective surgery</li> <li>• Clinically unstable patients</li> <li>• Inability of giving valid informed consent</li> </ul>
<b>Variables under study:</b>	<p><b>Patient-related:</b></p> <ul style="list-style-type: none"> <li>• Sex (Male/Female)</li> <li>• Age (years)</li> <li>• BMI (kg/m<sup>2</sup>)</li> <li>• aCCI (age-adjusted Charlson comorbidity index)</li> <li>• Previous abdominal surgery (yes/no)</li> </ul> <p><b>Disease-related:</b></p> <ul style="list-style-type: none"> <li>• Acute diverticulitis / Acute cholecystitis / Obstructed Hernia / Other (specify)</li> <li>• Severity grading of diverticulitis according to Hinchey classification (1a / 1b / 2 / 3 / 4) (if applicable)</li> <li>• Severity grading of cholecystitis according to Tokyo guidelines (I / II / III)(if applicable)</li> <li>• Site of hernia (if applicable)</li> </ul>

**Treatment-related:**

- Amount of robotic surgeries performed in the institution
- Expertise of surgeon (number of robotic surgeries performed)
- Type of procedure performed
- Time of intervention (hh:mm AM/PM)
- Operative time (min)
- Intraoperative complications (Bleeding or damage to major vessels / Organ injuries requiring reconstruction or resection / Unexpected medical conditions interrupting or changing the planned procedure)
- Conversion (yes/no)
- Reasons for conversion (bleeding / organ damage / adhesions or technical difficulties / anaesthesiologic contraindications)
- Drain placement (yes/no)

**Recovery-related:**

- ICU (yes/no)
- Clavien-Dindo (I/II/IIIa/IIIb/IVa/IVb/V)
- Type of complications
- Treatment of complications
- Death (yes/no)
- Time to first flatus (postoperative day)
- Time to first mobilization (postoperative day)
- Time to oral feeding (postoperative day)
- Length of stay (days)
- 90 days mortality
- 90 days readmission

**Cost analysis-related:**

- Number of consultant surgeons (specifying whether ordinary or dedicated)
- Number of non-consultant surgical assistants (specifying whether ordinary or dedicated)
- Number of anaesthetic consultant (specifying whether ordinary or dedicated)
- Number of non-consultant anaesthetic (specifying whether ordinary or dedicated)
- Number of theatre nurse scrubber (specifying whether ordinary or dedicated)
- Number of theatre nurse circulating (specifying whether ordinary or dedicated)
- Porter staff (specifying whether ordinary or dedicated)
- Amount of the following surgical instruments used: Maryland bipolar forceps, Fenestrated bipolar forceps, Permanent cautery hook, Cadiere forceps, Hot shears, Prograsp forceps, Vessel sealer, Harmonic Ace, staplers with reloads, sutures, drains, hemostatic consumables, diathermy consumables, scrub suits, dressings, drapes.

<b>Follow-up</b>	90 days
<b>Statistical methods, Propensity Score Matching, Sample size</b>	<p>Data will be expressed as median and interquartile range (IQR) and number and relative percentage. Normal distribution of continuous variables will be assessed with the Kolmogorov-Smirnov test.</p> <p>Continuous variables will be analyzed using the student t-test or Mann-Whitney test and categorical variables using Fisher exact test or Chi-Square test as appropriate.</p> <p>Significant variables (<math>p &lt; 0.05</math>) at univariate and well-known variables affecting outcomes will be used to run the matching.</p> <p>All statistics will be 2-tailed and statistical significance will be accepted when <math>p &lt; 0.05</math>. All statistical analyses will be performed using IBM SPSS Statistics 27.</p>
<b>Duration of the Study</b>	2023 - Ongoing
<b>Ethical Committee</b>	Comitato Etico Università Federico II -A.O.R.N. Cardarelli
<b>Dataset and Datadictionary</b>	Dataset and Datadictionary will be provided to all participant centres.
<b>Data management</b>	University of Naples “Federico II” will be responsible for collecting case report forms, controlling the quality of the reported data and generating reports and analyses, in cooperation with the Study Coordinator.
<b>Insurance</b>	NA
<b>References</b>	<ol style="list-style-type: none"> <li>1. Jung M, Morel P, Buehler L, Buchs NC, Hagen ME. Robotic general surgery: current practice, evidence, and perspective. <i>Langenbecks Arch Surg.</i> 2015;400:283–92. <a href="https://doi.org/10.1007/s00423-015-1278-y">https://doi.org/10.1007/s00423-015-1278-y</a>.</li> </ol>

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