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1 **TYPE OF ARTICLE:** Letter to Editors

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3 **TITLE:** Respiratory patterns in older patients following cardiac surgery: Trial of the
4 novel 'respi-R8' monitor

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25 surgery: trial of the novel 'respi-R8' monitor.

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28 submission.

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31 **TITLE:** Respiratory patterns in older patients following cardiac surgery: Trial of the
32 novel 'respi-R8' monitor

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34 **To the Editors,**

35 The respiratory rate (RR) is an important vital sign – useful to many specialties
36 including respiratory, anaesthetics and cardiology. Little is known about the
37 postoperative RR patterns in elderly patients after cardiac surgery. Therefore, this
38 preliminary study assessed RR continuously in older patients directly after tracheal
39 extubation following cardiac surgery using a non-invasive monitoring device that
40 measures the humidity of exhaled air, the respiR8® (Anaxsys). The respiR8
41 measures the RR independently from chest-wall movements; therefore, it can detect
42 both central (e.g. opioid-induced) and peripheral (e.g. obstructive) apnoea. We also
43 compared the results obtained with the respiR8® monitor with the ECG-derived RR
44 displayed on a standard recovery room monitor.

45 Twelve consecutive patients over the age of 60 years undergoing elective cardiac
46 surgery were monitored in August and September 2014 at King's College Hospital,
47 London. Their average age was 70.7 ± 9 years and average body mass index was
48 25.3 ± 4.6 ; one patient had a history of chronic obstructive pulmonary disease and
49 eight had histories of obstructive sleep apnoea; there were six non-smokers and two
50 active smokers. Due to the small sample size it was not possible to control for these
51 factors. It was a pilot study, intended to give results of interest in the planning of
52 larger studies.

53 They underwent the following procedures: coronary artery bypass graft (CABG) in
54 three patients, combined CABG and valve replacement in five, valve replacement in
55 three, and closure of an atrial septal defect in one. All operations were performed on
56 cardiopulmonary bypass with standard anaesthetic techniques. Postoperatively, all
57 patients were transferred to the cardiac recovery unit, where they were ventilated for
58 up to 6 hours. Once weaned off the ventilator, the patients were given an oxygen
59 mask with the respiR8® device attached. Patients were assessed using the respiR8®
60 device within 1 hour of extubation. The respiratory rate using both the respiR8®
61 device and an electrocardiography (ECG) monitor and oxygen saturation using pulse
62 oximetry were recorded every 10 min during the first hour of monitoring. Standard

63 postoperative care was maintained throughout the monitoring period. Each patient
64 received supplemental oxygen via a face mask. Routine observations and
65 investigations were performed, including regular arterial blood sampling.

66 Analgesia was maintained with continuous morphine infusion at a dose of 0.5–2
67 mg/h and regular intravenous paracetamol (1 g). No other drug with a significant
68 impact on breathing pattern was administered during this period. Cardiovascular
69 stability was maintained in all patients as per standardised protocols.

70 Apnoea was defined as an RR \leq 6 breaths/min for \geq 10 s [1], tachypnoea as an RR \geq
71 18 breaths/min, and bradypnoea as an RR \leq 10 breaths/min. Tachypnoea,
72 bradypnoea, and apnoea were referred to as 'abnormal respiratory events' (AREs).

73 Within the first hour after extubation, each patient had at least four AREs: 8 had a
74 mean of 5.1 episodes of bradypnoea, 9 had a mean of 2.9 episodes of tachypnoea,
75 and 8 had a mean of 2.5 episodes of apnoea. Seven patients experienced all three
76 AREs.

77 All patients were haemodynamically stable during the observation period with normal
78 mean arterial pressure, adequate filling pressure, and normal heart rate. All patients
79 received a mean dose of 1.3 mg/h of morphine in order to provide analgesia. The
80 arterial blood gas results assessed during the observation period revealed that two
81 patients had a lower than normal pH.

82 There was a difference in the clinical information gathered by continuous as opposed
83 to intermittent monitoring. Abnormal patterns were more likely to be identified by
84 continuous monitoring when compared with intermittent RR monitoring. Out of a
85 possible 72 opportunities (six per patient), the ECG failed to provide any reading at
86 all on 27 occasions. The ECG-derived RR monitoring showed larger negative
87 deviations, that is, lower RRs.

88 All of our patients experienced great variability in RRs and AREs during the 1-hour
89 observation period after tracheal extubation. All patients had just undergone major
90 cardiac surgery and were older than 60 years of age. There are many possible
91 reasons for AREs, including surgical stress, post-reperfusion syndrome, pain,
92 agitation, metabolic acidosis and continuous opioid infusions. The clinical
93 significance of these unexpected results warrants further investigation.

94 Our findings concurred with recent studies. For example, Smith *et al.* identified a
95 group of patients whose RRs were erratic over the measurement period. These
96 patients showed greater variability than healthy volunteers [2].

97 In our study, the respiR8® was more sensitive at detecting AREs than ECG-derived
98 RR monitoring. In a similar study comparing ECG-derived RR monitoring with
99 respiR8® monitoring, ECG-derived readings were prone to artefacts. Common
100 movements such as coughing and shivering led to bias in the results [3]. ECG-
101 derived RRs were more likely to fail to give any reading at all. When readings were
102 available, they were likely to differ from those given by the respiR8® monitor. In this
103 observational case series, oxygen saturation monitors proved unreliable - on seven
104 occasions where patients had at least one episode of apnoea, their oxygen
105 saturation levels were recorded as 94% or above, similar to our study.

106 While ECG-derived RR monitoring has been shown to be a weak indicator of
107 respiratory dysfunction, we found that the respiR8® was reliable and easy to use.

108 Our study was a preliminary study and had a number of limitations. The study did not
109 control for confounding factors such as previous history of respiratory compromise or
110 surgical technique. The study did not deal with any suggested interventions in
111 order to normalise RRs. In addition, postoperative RR-related complications, for
112 example, severe pain or pneumothorax, were not recorded. Postoperative outcome
113 variables, such as postoperative delirium, and other organ morbidity variables were
114 not assessed either.

115 In conclusion, our preliminary study showed a high incidence of AREs 1 hour after
116 cardiac surgery in elderly patients. The continuous respiR8® monitor was more
117 sensitive than a standard ECG-derived RR monitor. Abnormal patterns were more
118 likely to be identified by continuous monitoring when compared with intermittent RR
119 monitoring. Given the small sample size, the results cannot be generalised.
120 However, they do provide helpful preliminary data for the design of larger
121 observational RR-related clinical outcome studies.

122

123 **Keywords:** respiratory rate, post-operative, respi-R8

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126 **CONFLICT OF INTEREST**

127 None of the authors have a conflict of interest to declare.

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129 **AUTHOR'S CONTRIBUTIONS**

130 Dr. Rachel Kathleen Frances McNulty

131 Group1 - Conception and design, Acquisition of data, Analysis and interpretation of
132 data

133 Group 2 - Drafting the article, Critical revision of the article

134 Group 3 - Final approval of the version to be published

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148 Group 3 - Final approval of the version to be published

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153

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