Breathing variability—implications for anaesthesiology and intensive care

Oscar F. C. van den Bosch*©, Ricardo Alvarez-Jimenez, Harm-Jan de Grooth, Armand R. J. Girbes and Stephan A. Loer

Abstract
The respiratory system reacts instantaneously to intrinsic and extrinsic inputs. This adaptability results in significant fluctuations in breathing parameters, such as respiratory rate, tidal volume, and inspiratory flow profiles. Breathing variability is influenced by several conditions, including sleep, various pulmonary diseases, hypoxia, and anxiety disorders. Recent studies have suggested that weaning failure during mechanical ventilation may be predicted by low respiratory variability. This review describes methods for quantifying breathing variability, summarises the conditions and comorbidities that affect breathing variability, and discusses the potential implications of breathing variability for anaesthesia and intensive care.

Keywords: Respiratory variability, Control of breathing, Spontaneous respiration, Coefficient of variation, Detrended fluctuation analysis, Entropy analysis, Perioperative period, Spontaneous breathing trial

Background
The control of breathing involves a complex system that balances the opposing goals of efficiency, redundancy, responsiveness, and stability [1]. It is characterised by myriad inputs, internal pacemakers, positive and negative feedback loops, and nonlinear interactions between different components (Fig. 1). This results in fluctuations in breathing parameters, including the respiratory rate, tidal volume, and airflow profiles (Table 1).

If this regulation is too rigid, respiratory variability is low or absent, and the respiratory system cannot adequately react to stimuli. In contrast, if the respiratory system is overreacting to internal and external stimuli, the system shows large fluctuations and loses control.

The exact determinants of breathing variability are not precisely known; however, it has been shown that the increase and decrease in breathing variability are strongly associated with pathological states. Normal breathing variability is influenced by several factors, such as aging, cognitive load, sleep pattern, and hypoxia, as well as medical conditions such as anxiety, obstructive or restrictive lung disease, and arterial hypertension. During anaesthesia and intensive care, additional factors, such as drugs and the effects of mechanical ventilation, may also influence breathing variability.

This review describes the methods for quantifying breathing variability, summarises the conditions and comorbidities that affect breathing variability, and discusses the potential implications of breathing variability for anaesthesia and intensive care.

Normal respiratory variability
The regulation of breathing facilitates adequate gas exchange for metabolic needs. The impulses for inspiration and expiration are generated within the respiratory centre in the medulla oblongata after receiving and processing input from various subsystems (Fig. 1 and Table 2). Consequently, respiration is characterised by constant fluctuations in rate, rhythm, depth, and duration [2, 3]. For instance, this physiological variability...
may range between 19 and 34% for tidal volume and between 16 and 22% for respiratory rate, expressed as the coefficient of variation, in awake persons (Table 3).

The respiratory system receives input mainly from central and peripheral chemoreceptors, mechanoreceptors within the airways and alveoli, locomotion receptors of muscles and joints, and the (para)limbic system. The breathing pattern may be influenced to an extent and voluntarily controlled, shortly, via input from the cerebral cortex. For this purpose, the forebrain sends signals to the respiratory centre via independent pathways, over-ruling other inputs. Further factors include afferent input from the vagus nerve and its branches, such as the superior laryngeal nerves. External vagal stimulation has been shown to decrease respiratory rate [4], while stimulation of the superior laryngeal nerves may affect the chest wall and airway muscles [5]. Inspiratory time and tidal volume remain strongly correlated, suggesting a constant flow at a steady chemical drive [6, 7].

Different models have been developed to predict respiratory variability. One of the first models used a semi-mechanistic approach based on feedback loops of
measured (cardiac output and mixed venous blood CO₂ partial pressure) and estimated parameters (CO₂ sensitivity, mean lung volume for CO₂, circulating time) and integrated the result into a compartmental model using differential Eqs. [8]. For these calculations, several assumptions were necessary, including constant hemodynamic parameters, absence of intracardiac or pulmonary shunting, and instant intra-alveolar equilibration of CO₂ and oxygen tensions. Another more recent approach used spectral analysis of all variables to predict oscillatory rhythmicity [9]. This model incorporated inspiratory and expiratory times and volumes, as well as end-tidal CO₂ partial pressures and driving parameters. Of note, both models were based on measurements of healthy participants and can be extrapolated to patients with caution.

Quantification of respiratory variability

Different methods are available for the quantification of breathing variability, including quantitative time series analysis, detrended fluctuation analysis, entropy analysis, frequency distribution analysis, spectrum analysis, and power-law analysis [1]. In this section, we focus on the first three techniques.

Quantitative time series analysis

A quantitative time series analysis evaluates the standard deviation or interquartile range over time, such as the standard deviation of tidal volume or respiratory rate. The coefficient of variation (CV), defined as the ratio of the standard deviation to the mean, shows the extent of variability in relation to the mean of the data series (Eq. 1). The coefficient of variation is useful for comparing datasets with different units or means. It can be used as a measure of short- and long-term variations, depending on the subsets of data analysed. The extent of variability between successive breaths was calculated as the root mean square of successive differences (RMSSD) over consecutive breaths (short-term variability, Eq. 2). A quantitative time series analysis shows the overall degree of variability or “quantitative variability”.

\[
CV(x) = \sqrt{\frac{1}{N} \sum_{i=1}^{N} (x_i - \bar{x})^2}
\]  
\[\text{RMSSD}(x) = \sqrt{\frac{1}{N} \sum_{i=1}^{N} |x_{i-1} - x_i|^2}
\]  

Quantitative time series analysis: CV: coefficient of variation; RMSSD: root mean square of successive differences; N: number of samples; x: measured variable; \(\bar{x}\): mean of x.

Detrended fluctuation analysis

While quantitative time series analysis is used to measure short- and long-term variations, detrended fluctuation analysis (DFA) is used to detect long-range correlations in time series [10, 11]. We will address detrended fluctuation analysis as “correlated variability.” It is based on the assumption that variations are due to extrinsic stimuli that cause local effects or the intrinsic dynamics of the system causing long-range correlations. To quantify the intrinsic variability of the system, the local effects are subtracted. The algorithm consists of several steps, starting with a time series of the measured variable \(x\), such as tidal volume or respiratory rate. A new integrated time series \(X_T\) is calculated by summing the differences between the individual value \(x_i\) and the average \(\bar{x}\) for all values (Eq. 3). Subsequently, this trend function \(X_T\) is divided into epochs of length \(n\) and the local least squares fit (local trend) \(Y_T\) within this time window is subtracted. The fluctuation \(F(n)\) is then calculated as the root mean square of the integrated and detrended time series (Eq. 4). Finally, this process is repeated over different epochs \(n\) and a log–log graph of \(F(n)\) against \(n\) is constructed. The slope of a straight line fit yields the parameter \(\alpha\), which characterises long-term correlations (Eq. 5). While a higher \(\alpha\)-value of 0.5 indicates a time series without any long-term correlations, an increased \(\alpha\)-value would suggest the presence of such correlations. One major disadvantage of DFA is that it requires large
| Factor               | Reference                  | Respiratory measurements | Measure of variability |
|----------------------|----------------------------|--------------------------|------------------------|
| **Age**              | Tobin et al. [26]          | Inductive plethysmography| CV                     |
|                      |                            |                          | Young versus old subjects: |
|                      |                            |                          | RR: 0.16 versus 0.17   |
|                      |                            |                          | TV: 0.22 versus 0.28*   |
|                      |                            |                          | MV: 0.22 versus 0.27*   |
|                      |                            |                          | IT: 0.19 versus 0.21    |
| Peng et al. [11]     | Inductive plethysmography  | DFA (α)                  | Healthy young versus healthy elderly subjects (male): |
|                      |                            |                          | BT: 0.68 ± 0.07 versus 0.66 ± 0.08* |
|                      |                            |                          | Healthy young versus healthy elderly subjects (female): |
|                      |                            |                          | BT: 0.70 ± 0.07 versus 0.67 ± 0.06 |
| **Sleep**            | Hudgel et al. [16]         | Nasal CPAP mask, pneumotachography | CV                     |
|                      |                            |                          | Awake versus NREM sleep (elderly): |
|                      |                            |                          | TV: 0.24 ± 0.04 versus 0.25 ± 0.04 |
|                      |                            |                          | MV: 0.20 ± 0.02 versus 0.25 ± 0.05 |
|                      |                            |                          | IT: 0.15 ± 0.02 versus 0.11 ± 0.01 |
|                      |                            |                          | BT: 0.15 ± 0.02 versus 0.10 ± 0.05* |
|                      |                            |                          | Awake versus NREM sleep (young): |
|                      |                            |                          | TV: 0.23 ± 0.04 versus 0.11 ± 0.02* |
|                      |                            |                          | MV: 0.14 ± 0.02 versus 0.10 ± 0.11 |
|                      |                            |                          | IT: 0.17 ± 0.02 versus 0.08 ± 0.01* |
|                      |                            |                          | BT: 0.16 ± 0.02 versus 0.08 ± 0.01* |
| Rostig et al. [17]   | Full-face mask, pneumotachography | CV                    | NREM versus REM sleep: |
|                      |                            |                          | TV: 0.11 ± 0.01 versus 0.28 ± 0.08* |
|                      |                            |                          | MV: 0.11 ± 0.03 versus 0.21 ± 0.09* |
|                      |                            |                          | RR: 0.09 ± 0.03 versus 0.20 ± 0.04* |
|                      |                            |                          | IT: 0.10 ± 0.02 versus 0.15 ± 0.03 |
|                      |                            |                          | DFA, STC (α1)           |
|                      |                            |                          | TV: 0.89 ± 0.09 versus 1.18 ± 0.14 |
|                      |                            |                          | MV: 0.95 ± 0.10 versus 1.18 ± 0.16 |
|                      |                            |                          | RR: 0.79 ± 0.06 versus 0.95 ± 0.14 |
|                      |                            |                          | IT: 0.68 ± 0.04 versus 0.76 ± 0.10 |
|                      |                            |                          | DFA, LTC (α2)           |
|                      |                            |                          | TV: 0.51 ± 0.09 versus 0.81 ± 0.15 |
|                      |                            |                          | MV: 0.51 ± 0.09 versus 0.77 ± 0.13 |
|                      |                            |                          | RR: 0.57 ± 0.05 versus 0.85 ± 0.12 |
|                      |                            |                          | IT: 0.55 ± 0.07 versus 0.76 ± 0.11 |
| **Hypertension**     | Anderson et al. [30]       | Inductive plethysmography| RMSSD                  |
|                      |                            |                          | Lower versus upper tertile of blood pressure: |
|                      |                            |                          | TV: 147 ± 38 versus 219 ± 22* |
|                      |                            |                          | MV: 1.84 ± 0.32 versus 1.91 ± 0.46* |
|                      |                            |                          | RR: 3.56 ± 0.50 versus 4.72 ± 0.73* |
| **Children with anxiety disorder** | Pine et al. [32] | Spirometry and respiratory canopy | SD                     |
|                      |                            |                          | Anxiety disorder versus normal controls: |
|                      |                            |                          | TV: 98 ± 101 versus 39 ± 27* |
|                      |                            |                          | MV: 2.0 ± 2.0 versus 0.8 ± 0.4* |
|                      |                            |                          | RR: data not reported, P = 0.06 |
| **Panic disorder**   | Martinez et al. [34]       | Spirometry and respiratory canopy | SD                     |
|                      |                            |                          | Panic disorder versus normal controls: |
|                      |                            |                          | TV: 191 ± 184 versus 84 ± 43* |
|                      |                            |                          | MV: 2.36 ± 1.80 versus 1.37 ± 0.68* |
|                      |                            |                          | RR: 4.59 ± 2.75 versus 3.58 ± 2.43 |
|                      |                            |                          | MSSD                   |
|                      |                            |                          | TV: 308 ± 358 versus 144 ± 152* |
|                      |                            |                          | MV: 4.27 ± 5.05 versus 2.12 ± 1.84* |
|                      |                            |                          | RR: 6.90 ± 7.75 versus 5.38 ± 5.13 |
| Factor             | Reference                  | Respiratory measurements          | Measure of variability                                      |
|--------------------|----------------------------|-----------------------------------|------------------------------------------------------------|
|                    |                            | Inductance plethysmography CV     | Panic disorder versus normal controls:                  |
|                    |                            |                                   | TV: 0.54 ± 0.22 versus 0.33 ± 0.15 (standing)*            |
|                    |                            |                                   | RR: 0.39 ± 0.17 versus 0.45 ± 0.10 (standing)*           |
|                    |                            |                                   | TV: 0.23 ± 0.10 versus 0.32 ± 0.27 (supine)             |
|                    |                            |                                   | RR: 0.29 ± 0.12 versus 0.32 ± 0.11                     |
|                    |                            | LLE                               | Panic disorder versus normal controls:                   |
|                    |                            |                                   | 0.10 ± 0.01 versus 0.086 ± 0.02 (standing)*             |
|                    |                            |                                   | 0.09 ± 0.02 versus 0.09 ± 0.02 (supine)                 |
|                    |                            | ApEn                              | Panic disorder versus normal controls:                   |
|                    |                            |                                   | 0.40 ± 0.13 versus 0.27 ± 0.12 (standing)*              |
|                    |                            |                                   | 0.30 ± 0.00 versus 0.29 ± 0.12 (supine)                 |
| Cognitive load     | Vlemincx et al. [21]       | Inductance plethysmography CV     | Complex arithmetic task versus baseline:                |
|                    |                            |                                   | TV: 0.36 ± 0.16 versus 0.24 ± 0.14*                     |
|                    |                            |                                   | MV: 0.26 ± 0.08 versus 0.20 ± 0.06*                     |
|                    |                            |                                   | RR: 0.18 ± 0.08 versus 0.16 ± 0.07                     |
|                    |                            |                                   | AR Complex arithmetic task versus baseline:             |
|                    |                            |                                   | TV: 0.11 ± 0.23 versus 0.20 ± 0.18*                     |
|                    |                            |                                   | MV: 0.30 ± 0.24 versus 0.28 ± 0.18                     |
|                    |                            |                                   | RR: 0.12 ± 0.15 versus 0.26 ± 0.20*                    |
|                    |                            | Grassmann et al. [22] Nasal capnometry CV | High demanding mental multitask versus baseline:  |
|                    |                            |                                   | RR: 0.13 ± 0.05 versus 0.19 ± 0.09*                     |
| COPD               | Loveridge et al. [37]      | Inductance plethysmography CV     | COPD versus normal controls:                            |
|                    |                            |                                   | TV: 0.253 versus 0.337*                                 |
|                    |                            |                                   | MV: 0.221 versus 0.280*                                 |
|                    |                            |                                   | RR: 0.170 versus 0.220                                  |
|                    |                            |                                   | IT: 0.178 versus 0.229                                  |
| Asthma             | Hmeidi et al. [38]         | Structured light plethysmography IQR | Asthma (prebronchodilator) versus normal controls:     |
|                    |                            |                                   | RR: 3.93 (2.57) versus 3.32 (2.2)                      |
|                    |                            |                                   | IE50: 0.63 (0.32) versus 0.47 (0.18)*                   |
|                    |                            |                                   | Asthma (prebronchodilator) versus asthma (postbroncho dilator): |
|                    |                            |                                   | RR: 3.93 (2.57) versus 4.62 (2.34)                      |
|                    |                            |                                   | IE50: 0.60 (0.32) versus 0.60 (0.38)*                   |
| Asthma             | Seppa et al. [39]          | Impedance pneumography CSRmin     | High-risk group versus low-risk group:                  |
|                    |                            |                                   | Flow-volume curve 0.995 [0.984–0.999] versus 0.998 [0.994–0.999]* |
|                    |                            |                                   | NLmin High-risk group versus low-risk group:            |
|                    |                            |                                   | Flow signal 14.3 [0.00–48.7] versus 30.3 [0.00–42.7]* |
| Factor                          | Reference                        | Respiratory measurements       | Measure of variability                                                                 |
|--------------------------------|----------------------------------|--------------------------------|----------------------------------------------------------------------------------------|
| Restrictive lung disease       | Brack et al. [42]                | Inductance plethysmography CV  | CV restrictive lung disease versus normal controls: TV 0.22 ± 0.05 versus 0.50 ± 0.20*  |
|                                |                                  |                                | IT 0.22 ± 0.05 versus 0.33 ± 0.12*                                                    |
|                                |                                  |                                | ET 0.22 ± 0.07 versus 0.41 ± 0.19*                                                    |
|                                |                                  |                                | MV 0.24 ± 0.06 versus 0.42 ± 0.16*                                                    |
|                                |                                  |                                | AR TV 0.42 ± 0.14 versus 0.23 ± 0.12*                                                 |
|                                |                                  |                                | IT 0.25 ± 0.17 versus 0.21 ± 0.14                                                    |
|                                |                                  |                                | ET 0.28 ± 0.13 versus 0.12 ± 0.15*                                                   |
|                                |                                  |                                | MV 0.39 ± 0.16 versus 0.27 ± 0.14                                                    |
| Lung dysmaturity in infancy    | Fouzas et al. [43]               | Full face mask, flowmeter CV   | CV preterm non-CLDI infants versus term infants: TV 0.09 ± 0.03 versus 0.09 ± 0.02     |
|                                |                                  |                                | Preterm moderate/severe CLDI versus term infants: TV 0.07 ± 0.02 versus 0.09 ± 0.02*  |
|                                |                                  |                                | 2D dispersion, Poincaré TV 147 ± 82.2 versus 143 ± 64.3                               |
|                                |                                  |                                | Preterm moderate/severe CLDI versus term infants: TV 147 ± 82.2 versus 143 ± 64.3*    |
|                                |                                  |                                | TV 58.9 ± 40.7 versus 143 ± 64.3                                                     |
|                                |                                  |                                | 3D dispersion, Poincaré TV 1156 ± 906 versus 1073 ± 670                               |
|                                |                                  |                                | Preterm moderate/severe CLDI versus term infants: TV 1156 ± 906 versus 1073 ± 670*    |
|                                | Usemann et al. [44]              | Full face mask, flowmeter CV   | CV preterm versus term infants: Rint 20.2 ± 8.4 versus 29.6 ± 14.9                    |
| After major abdominal surgery  | Van den Bosch et al. [56]        | Impedance pneumography CV      | CV RR versus TV 0.21 ± 0.06 versus 0.37 ± 0.12*                                        |
|                                |                                  |                                | RR versus TV 0.21 ± 0.06 versus 0.37 ± 0.12*                                          |
|                                | Bradley et al. [63]              | Capnography CV                 | CV before versus after sedation interruption, low MODS: RR 0.17 ± 0.08 versus 0.28 ± 0.16* |
| Organ dysfunction syndrome in  |                                  |                                | RR before versus after sedation interruption, high MODS: RR 0.23 ± 0.12 versus 0.20 ± 0.12 |
| ICU                            |                                  |                                | RMSSD before versus after sedation interruption, low MODS: RR 0.86 ± 0.53 versus 1.7 ± 1.3 |
|                                |                                  |                                | RR before versus after sedation interruption, high MODS: RR 1.3 ± 0.79 versus 1.0 ± 0.79* |
|                                |                                  |                                | ApEn before versus after sedation interruption, low MODS: RR 0.48 ± 0.16 versus 0.46 ± 0.15 |
|                                |                                  |                                | RR before versus after sedation interruption, high MODS: RR 0.42 ± 0.16 versus 0.49 ± 0.17* |
|                                |                                  |                                | DFA, STC (α1) before versus after sedation interruption, low MODS: RR 0.64 ± 0.19 versus 0.70 ± 0.15 |
|                                |                                  |                                | RR before versus after sedation interruption, high MODS: RR 0.69 ± 0.09 versus 0.68 ± 0.17 |
For this purpose, the entropy analysis algorithm evaluates whether a sequence of data points of length \( m \) is similar to other sequences in the data within a specified tolerance \( r \). Subsequently, the difference between the logarithmic frequencies of similar runs of length \( m \) and \( m + 1 \) is measured. When the data contain several repetitive patterns, the approximate entropy (ApEn) is low; otherwise, the algorithm yields a higher ApEn. One shortcoming of the calculation of ApEn is its dependency on the choice of sequence length \( m \) and tolerance \( r \) because other choices of \( m \) and \( r \) can lead to different conclusions on the randomness of the data.

**Approximate entropy analysis:** \( r \): similarity criterion; \( n_{im}(r) \): number of patterns that are similar within \( r \); \( m \): pattern length; \( S_N \): sequence of \( N \) measurements; \( C_{im}(r) \): mean of all \( C_{im}(r) \) values; ApEn: approximate entropy.

\[
C_{im}(r) = \frac{n_{im}(r)}{N - m + 1} \tag{6}
\]

\[
\text{ApEn}(S_N, m, r) = \ln \frac{C_{im}(r)}{C_{m+1}(r)} \tag{7}
\]

**Conditions and diseases influencing respiratory variability**

Breathing variability is modified and influenced by several conditions, such as sleep, cognitive function, age, hypoxia, and diseases (Table 3).

### Sleep

In healthy participants, quantitative respiratory variability decreases during non-rapid eye movement (non-REM) sleep [15, 16]. In addition, breath-to-breath components display a strong relationship between one breath and another at a time lag of a few breaths (short-term correlations). The regulation of respiratory timing and drive is characterised by additional long-term correlations only during the transition from non-REM to REM sleep [17]. The quantitative variability of respiratory rate is augmented during REM sleep, compared to non-REM sleep, but less prominent than during wakefulness [18]. The decreased quantitative variability of respiration during non-REM sleep is a result of an autoregressive process (i.e. breaths depend linearly on their previous values), as well as a result of periodic oscillations and uncorrelated white noise [19]. The observed breath-to-breath dependence (autoregression) is likely caused by the central respiratory pattern generator, whereas periodic oscillations are more likely to originate from chemical feedback systems [19]. It is unknown whether similar changes can be observed during sedation and anaesthesia with preserved spontaneous respiration.

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**Table 3** (continued)

| Factor | Reference | Respiratory measurements | Measure of variability |
|--------|-----------|--------------------------|------------------------|
|        |           | DFA, LTC (\( q_2 \))     | Before versus after sedation interruption, low MODS: |
|        |           |                          | RR 1.17 ± 0.78 versus 0.33 ± 0.41 |
|        |           |                          | Before versus after sedation interruption, high MODS: |
|        |           |                          | RR 0.25 ± 0.22 versus 0.40 ± 0.35* |

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\( CV \): coefficient of variation; \( TV \): tidal volume; \( MV \): minute ventilation; \( IT \): inspiratory time; \( BT \): breath time; \( RR \): respiratory rate; \( DFA \): detrended fluctuation analysis; \( LTC \): long-term correlations; \( ApEn \): approximate entropy; \( AR \): autocorrelation at one breath lag; \( COPD \): chronic obstructive pulmonary disease; \( CLDI \): chronic lung disease of infancy (a measure of chaos); \( ApEn \): approximate entropy (a measure of regularity); \( AR \): autocorrelation; \( MODS \): multiple organ dysfunction syndrome.* 

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\[ X(t) = \sum_{i=1}^{t} |x(t) - \bar{x}| \tag{3} \]

\[ F(n) = \sqrt{\frac{1}{N} \sum_{t=1}^{N} [X(t) - Y(t)]^2} \tag{4} \]

\[ F(n) \propto n^\alpha \tag{5} \]

**Detrended fluctuation analysis:** \( x(t) \): measured variable; \( \bar{x} \): mean of \( x \); \( n \): epoch length; \( Y(t) \): local least-squares fit; \( N \): number of measurements; \( \alpha \): scaling component characterising the extent of long-term correlations.

**Entropy analysis**

Entropy analysis measures the degree of disorder or randomness in the data. In other words, it quantifies the amount of “surprise” or “new information” introduced to an otherwise predictable system. Entropy analysis, therefore, reflects the degree of complexity or “informational variability.” This elegant analysis approach is often applied to thermodynamics, but it can also be used to analyse breathing irregularities [13, 14]. Data are considered more irregular and unpredictable when observed patterns are not followed by similar patterns. For this purpose, the entropy analysis algorithm evaluates whether a sequence of data points of length \( m \) is similar to other sequences in the data within a specified tolerance \( r \). Subsequently, the difference between the logarithmic frequencies of similar runs of length \( m \) and length \( m + 1 \) is measured. When the data contain several repetitive patterns, the approximate entropy (ApEn) is low; otherwise, the algorithm yields a higher ApEn. One shortcoming of the calculation of ApEn is its dependency on the choice of sequence length \( m \) and tolerance \( r \) because other choices of \( m \) and \( r \) can lead to different conclusions on the randomness of the data.
**Cognitive load**
Changes in cognitive activity also influence respiratory variability [20]. Arithmetic tasks under stressful conditions significantly increased the quantitative variability of respiratory rate, tidal volume, and minute volume by 13%, 50%, and 30%, respectively, compared with measurements during restful watching of the documentary titled “The March of the Penguins” [21]. In another mental load experiment, however, quantitative and correlated variability of respiratory rate was reduced by 30% when volunteers were exposed to multiple tasks while assessing their perceptual speed, spatial orientation, and working memory capacity. Breathing variability was restored to its baseline values during the recovery period [22]. The duration and complexity of cognitive load have variable effects on respiratory variability.

**Exercise**
Increasing metabolic demands during physical exercise induces an increase in minute ventilation through the augmentation of the respiratory rate and tidal volume. However, the quantitative variability of the respiratory rate decreases during recumbent bicycle exercise [23]. The correlated variability also decreases, resulting in a more random pattern of breathing, which suggests that the respiratory control system may be operating in a less stable state during exercise [23]. The degree of athletic fitness altered the effect of exercise on breathing variability. In a cardiopulmonary exercise test at maximum oxygen consumption, sedentary volunteers showed a 40% higher quantitative variability of minute ventilation than professional football players from Brazil [24]. In patients with heart failure, a breathing pattern with cyclic fluctuations in minute ventilation during incremental exercise (“exertional oscillatory ventilation”) is a strong predictor of poor prognosis and reflects advanced disease [25].

**Age**
Respiratory variability increases with age. Healthy older adults [60–81 years] showed greater quantitative variability than younger participants [21–50 years], especially during non-REM sleep [16, 26]. Asleep older adults more often show an increase in upper airway resistance, which produces breathing fluctuations caused by mechanical limitations contributing to this breath-to-breath variability of tidal volume [16]. A study of long-term correlations (120 min) shows long-range correlations extending over hundreds of breathing cycles during sleep, and these correlations seem to degrade in older men [11]. The authors suggest that this loss of long-term correlations in breathing dynamics may be caused by intrinsic factors associated with aging such as neuronal dropout, loss of central signal integration, stiffening of the pulmonary parenchyma, and reduced chemoreceptor sensitivity. It is unknown whether these changes in quantitative and correlated variability of breathing are accompanied by changes in complexity or informational variability.

**Hypoxia and high altitude**
Specific changes in respiratory patterns are observed under hypoxic conditions [27, 28]. During the transition from normoxia to isocapnic hypoxia, the quantitative variability of minute ventilation increases by 70%, and that of tidal volume increases by 50%, without inducing periodic breathing [28]. This was mainly mediated by an increase in the random fraction, as the correlated variability decreased [28]. Hypoxic conditions at high altitudes induced similar changes. The incidence of periodic breathing with apneic episodes and their duration also increased as a function of altitude in non-acclimatized participants [29]. Breathing patterns oscillate periodically between clusters of breaths and periods of short apnea during NREM sleep under hypobaric hypoxia [27]. This pattern increases the variability of breathing, quantified in terms of the coefficient of variation. This periodic breathing pattern does not occur during wakefulness or REM sleep and cannot be observed during normocapnia.

**Arterial hypertension**
In addition, arterial hypertension may influence breathing variability. The quantitative variability of tidal volume, respiratory rate, and minute ventilation was significantly higher in a cohort of women at rest with elevated systolic blood pressure [30]. The incidence of apneic events (> 10 s) during resting wakefulness increased more than twofold in patients with elevated blood pressure, whereas the incidence of these breathing pauses did not differ during overnight sleep. It has not been established if periodic breathing is a cause or consequence of long-term hypertension; however, it has been hypothesised that increased sympathetic and/or decreased parasympathetic activity may account for the breathing pattern.

**Endotoxin**
The systemic inflammatory response to endotoxins induces alterations in respiratory frequency and minute ventilation, as well as the respiratory pattern [31]. Even though the overall quantitative variability remained unaltered, the correlated variability of respiratory rate was 42% higher after exposure to endotoxins than placebo [31]. These changes were functions of changes in $P_{CO_2}$, suggesting a role of chemical feedback loops. Interestingly, ibuprofen suppressed the increase in the correlated behaviour of respiratory frequency.
Anxiety
Marked differences in breathing variability have been reported in children and adolescents with anxiety disorders, such as panic disorder [32, 33]. During resting wakefulness, patients at risk of panic attacks have a twofold higher quantitative variability of tidal volume than healthy controls. This difference persisted even after treatment [34]. In patients with hyperventilation disorder, there is an augmented degree of quantitative variability and a concomitant augmentation in the complexity of tidal volume variability [35]. One study reported increased quantitative and informational respiratory variability in patients with panic disorder only in the standing (but not in the supine) position. The authors hypothesised that this difference may be attributed to a diminished vagal tone in patients with panic disorders [36].

Obstructive lung disease
Asthma and chronic obstructive pulmonary disease (COPD) affect breathing variability in different ways. Patients with COPD have higher quantitative variabilities of minute ventilation and tidal volume than age- and sex-matched controls [37]. They have significantly fewer sighs; even after the exclusion of sighs from analysis, respiratory variability was lower in patients with COPD [37]. This reduced variability in breathing patterns may reflect changes in the mechanics of the lung and chest wall or neural adjustments in breathing control.

In contrast, the variability of respiration is increased in patients with asthma, and this effect is related to disease severity. The quantitative variability of tidal breathing parameters in children with asthma aged 7–16 years was 25% greater than that in age-matched children, independent of the use of a bronchodilator [38]. In young patients [3–7 years] with risk factors for asthma, compared to a control group, the quantitative variability of tidal flow is significantly increased [39]. Patients treated with inhalational corticosteroids, compared to the low-risk group, showed a normal tidal flow pattern, suggesting that control medication modifies disease activity and lung function variability.

These distinctively different effects of COPD and asthma on breathing variability were confirmed by a study that used the forced oscillation technique [40] to determine the variability of respiratory impedance and other airway properties. In this study, the temporal dynamics of respiratory impedance was used to distinguish the asthma and COPD groups [41].

Restrictive lung disease
Variability in breathing was also reduced in patients with restrictive lung disease. For instance, during resting wakefulness, the quantitative variability of tidal volume was reduced by 56%, inspiratory time was reduced by 33%, expiratory time was reduced by 46%, and minute ventilation was reduced by 43% compared to those of healthy participants [42]. Concurrently, the correlated variability increased, suggesting that patients with restrictive lung disease adopt a constrained breathing pattern.

Lung prematurity
Variability in breathing is also influenced by lung maturity. In preterm infants (i.e. born before 37 gestational weeks), quantitative variability of tidal volume and end-tidal expiratory CO₂ is significantly lower in patients with chronic lung disease of infancy than in patients without supplementary oxygen or CPAP [43]. At the postmenstrual age of 42–50 weeks, the quantitative variability of airway resistance is significantly higher in term infants than in preterm infants [44]. In preterm born infants, lower quantitative variability of tidal volume at a postmenstrual age of 44 weeks is an important predictor of re-hospitalization due to respiratory disease in infancy [45].

Perioperative period
Various drugs administered during the perioperative period influence breathing control. Propofol decreases the respiratory response to hypoxia and hypercarbia [46, 47] resulting in decreased tidal volume and minute ventilation. Opiates cause dose-dependent hypoventilation mainly through a decrease in respiratory rate [48]. This respiratory depression is often accompanied by increased quantitative tidal volume variability [49]. Midazolam reduces minute ventilation mainly through decreases in tidal volume and, to a lesser extent, respiratory rate [50]. On the other hand, s-ketamine activates breathing with an increase in respiratory rate and inspiratory time and can antagonise opiate-induced hypoventilation [51–55]. The exact effects of these drugs on respiratory variability have not been established. The quantitative variability of respiratory rate was lower than that of tidal volume during the first 24 postoperative hours in patients undergoing major abdominal surgery. These findings suggest that the adaptations of alveolar ventilation to metabolic needs may be predominately achieved by variations in tidal volume [56].
Intensive care
During weaning from mechanical ventilation, variability in breathing may be valuable for clinical decision-making [57]. Weaning is often preceded by a spontaneous breathing trial (SBT) in which patients are disconnected from ventilatory support to assess the adequacy of their respiratory function. In patients recovering from a systemic inflammatory response syndrome, quantitative variability of tidal volume during a 30 min SBT was significantly lower in patients requiring reinstitution of non-invasive or invasive mechanical ventilation within 48 h [58]. Similar observations were reported in another study of ventilated patients who underwent a 60-min SBT with oxygen supplementation. Patients with a lower quantitative variability of tidal volume during SBT more often require ventilatory support after weaning from mechanical ventilation [59].

Interesting findings were reported in a study of patients requiring prolonged ventilation (>7 days) [60]. During SBT with CPAP, the quantitative variability of tidal volume, but not respiratory rate, was higher in patients with failed weaning. Concurrently, the informational variability was higher in patients with failed weaning, suggesting a less predictable breathing pattern [61]. An important limitation of this study is that the patients were supported with a mean of 12 ± 4.6 cm H2O of pressure support although the aim was to assess the intrinsic variability of a patient [62].

The quantitative variability of the respiratory rate increases significantly during the reduction or interruption of sedation with propofol, midazolam, or their combination [63]. This restoration of respiratory rate variability is greater in patients with lower multiple organ dysfunction scores (MODS) [63].

Various studies on the prognostic value of respiratory variability are presented in Table 4. These findings are interesting; however, we need more data to determine

| Setting | Reference | Respiratory measurements | Measure of variability |
|---------|-----------|--------------------------|------------------------|
| Preterm infants | Usemann et al. [45] | Full face mask, flowmeter | CV |
| ICU weaning trial with 5 cmH2O PSV plus 5 cmH2O PEEP | Bien et al. [58] | Ventilator | CV |
| ICU spontaneous breathing trial without ventilatory support | Wysocki et al. [59] | Ventilator | CV |
| ICU weaning trial, CPAP 5 cmH2O | El Khatib et al. [60] | Ventilator | CV |
| ICU weaning trial with 12 ± 4.6 cmH2O PSV plus 5 cmH2O PEEP | Engeron [61] | Ventilator | ApEn |

CV coefficient of variation, TV tidal volume, IQR interquartile range, OR odds ratio, ICU intensive care unit, PSV pressure support ventilation, PEEP positive end-expiratory pressure, BT breathing time, PIF peak inspiratory flow, AUC area under the curve, ApEn approximate entropy* P < 0.05
the exact role of variability analyses in guiding clinical decisions.

Future research
There is increasing attention for temporal variations of physiologic variables, such as heart rate variability [64], and the amount of available data on breathing variability is increasing. Further research should elucidate the correlation between respiratory variability during the postoperative period and clinically relevant outcomes, such as postoperative morbidity and mortality [65]. Large amounts of physiological data are generated during anaesthesia and intensive care. Technological advances in data analysis, smart learning techniques, and artificial intelligence can facilitate the determination of patients at risk [66–68]. Integrating complex data from multiple sources may lead to improved risk stratification. Recent advances allow us to monitor respiratory function in a continuous and noninvasive manner [50, 56, 69–82]. This is important as postoperative pulmonary complications remain a major disease burden [83–87].

Conclusions
The variability of respiration over time may be a promising tool for identifying patients at risk of pulmonary complications. The variability of respiration is complex and not fully understood yet. Measuring the variability of a single parameter, such as the respiratory rate, does not necessarily reflect the variability of the respiratory system as a whole. The overall variability of breathing is decreased by COPD, restrictive lung disease, chronic lung disease of infancy, non-REM sleep, and highly demanding cognitive tasks. In contrast, it is increased in older adults during the performance of complex arithmetic tasks during hypoxia and in patients with asthma, hypertension, or anxiety disorder. Further research is required to elucidate the full potential of respiratory variability in critical care and anaesthesiology.

Abbreviations
CV: Coefficient of variation; RMSSD: Root mean square of successive differences; DFA: Detrended fluctuation analysis; ApEn: Approximate entropy; REM: Rapid eye movement; COPD: Chronic obstructive pulmonary disease; CLDI: Chronic lung disease of infancy; CPAP: Continuous positive airway pressure; SBT: Spontaneous breathing trial.

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Authors’ contributions
OvdB is responsible for the conception and drafting of this manuscript. RAJ is responsible for the conception and drafting of this manuscript. HdG is responsible for the conception and drafting of this manuscript. AG is responsible for the conception and drafting of this manuscript. SL is responsible for the conception and drafting of this manuscript. All authors reviewed the final manuscript for important intellectual content. All authors read and approved the final manuscript.

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