## **In2itive e-Diary**

Next generation eCOA platform with integrated respiratory assessment

The In2itive™ e-Diary is a robust handheld medical device with a high resolution touchscreen for gathering eCOA data. Using the proven Vitalograph® Fleisch Pneumotachograph flowhead, the In2itive e-Diary provides accurate measurement of respiratory endpoints in clinic or home settings.



### **Features**

- · Large, responsive touchscreen
- Fleisch pneumotachograph flowhead for respiratory endpoints e.g. FEV1 and PEF
- · Customizable alerts, reminders, and workflows to fit study protocols
- Date/time stamps on all data and results
- · Cellular/Mobile option transmits results direct to Vitalograph web portal
- Automatic secure end-to-end encrypted data transmission
- Live spirometry quality feedback
- Subject management via web portal or PC software
- Integrated animated training module
- Password protected
- CE marked and FDA cleared



The device can host any ePRO, eCOA or sponsor questionnaire using text, VAS, images, tick box, radio buttons, etc.

Pre integrated & validated ePROs:

#### Respiratory - asthma

ACQ MAPI
AQLQ(S)12+ MAPI
AQ20 Prof P Jones

ACT Quality Metric

#### Respiratory - COPD

EXACT Evidera
EXACT E-RS Evidera
CAT GSK

#### **Health status**

EQ5D-3L Euroqol EQ5D-5L Euroqol



# **Technical Specification**

Product:	Vitalograph 2120 In2itive e-Diary
Dimensions:	186 mm x 81 mm x 48 mm net
Weight:	0.3 kg net
Communications:	Minimum USB 2.0 Integrated Cellular/Mobile (optional)
Power Supply:	Universal Input 100 -240V, 50-60 Hz or USB Power 5V/500mA (as supplied) 3.7V Lithium-ion polymer battery
Flow Detection Principal:	Fleisch type Pneumotachograph capable of giving linear signals throughout the entire physiological range.
Volume Detection:	Flow integration sampling @ 100Hz
Volume Accuracy:	±3% or ±0.05L of the reading (ISO 26782:2009).
Flow Measurement Range:	Max. flow rate ±14 L/s Min. flow rate ±0.02 L/s
PEF Accuracy:	±10% or ±10L/min of the reading (ISO 23747:2015)
Back Pressure:	Less than 0.1kPa/L/sec @ 14L/s (ATS/ERS 2019)
Performance Standards: (the device meets or exceeds)	ATS/ERS 2019, ISO 23747:2015 & ISO 26782:2009
Safety Standards:	EN 60601-1:2006+A1:2013 with US deviations
EMC Standards:	EN 60601-1-2:2015
Home Use Standards:	EN 60601-1-11:2015
QA/GMP Standards:	EN ISO 13485, FDA 21 CFR 820, CMDR SOR/98-282 & JPAL

