SICATJMT+

SICAT JMT⁺ JAW MOTION ANALYSIS SYSTEM

Technical data and instructions for use – English

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1 USER INFORMATION

1.1 MANUFACTURER INFORMATION

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1.2 CONFIGURATION OF THE INSTRUCTIONS FOR USE

The instructions for use for the SICAT JMT⁺ jaw motion analysis system consist of two parts:

- 1. SICAT JMT⁺ system technical data and instructions for use
- 2. SICAT JMT⁺ instructions for use of the user software

The part SICAT JMT⁺ system technical data and instructions for use primarily contains information concerning the technical data and operation of the SICAT JMT⁺ system, as well as information on its safe operation in combination with the patient accessories such as the **SICAT Fusion Bite**. Information concerning accessory components is limited to key safety and maintenance measures and/or hygiene measures.



The exact compliance with the information in both parts of the instructions for use is required for proper use.

1.3 USED SYMBOLS



Green highlighting around the side of the instructions for use designates new information concerning product safety.



Warning instructions designate a potential danger to the health and safety of users and/or patients. The instructions explain the type of danger and how it can be avoided.

Notices designate a potential danger that can cause damage to the device. The NOTICE instructions explain the type of danger and how it can be avoided.



CE0197 The CE mark with code number 0197 of the notified body TÜV Rheinland LGA Products GmbH confirms the conformity of the measurement system with guideline 93/42/EEC (Medical Devices Directive).



Symbol for manufacturer and date of manufacture



Type BF device corresponding to DIN EN 60601-1



USB port



This symbol shows that pursuant to the Directive on Waste Electrical and Electronic Devices (2012/19/EU) and national legislation, a product cannot be disposed of via the household waste, and must be disposed of separately within Europe.



Icon for HF-transmitter (Bluetooth interface)



Article number of the measurement system / accessory



Serial number of the measurement system



An accessory that is intended for one-off use on a single patient during a single treatment.



Refer to instructions for use.

2 AREA OF USE AND SAFETY

2.1 INTENDED USE

Based on the lower jaw movements of the patient, the SICAT JMT⁺ system calculates all the necessary parameters with the goal of a supporting diagnosis for the following indications:

- · supporting diagnosis for treating a diseased mandibular joint
- · supporting diagnosis for creating occlusal mouth-guards
- · supporting diagnosis for creating dentures
- · supporting diagnosis for aesthetically functional reconstruction with or without tooth implants
- · supporting diagnosis for treating craniomandibular dysfunction (CMD)

The measurement system also enables the exporting of the gathered data for further processing with CAD/CAM or PACS systems.

The measurement system may only be used by trained dentists. The area of use is limited to dental systems. A measurement is carried out within 15 minutes and should not be applied in the case of open wounds in the oral and head area; it can be applied to patients who are over 10 years old and who are mentally capable of exactly following the instructions of the operator.

2.1.1 USE

The SICAT JMT⁺ system is an electronic recording system that is based on 3D ultrasound measurements. SICAT JMT⁺ systems record the lower jaw movements of the patient in all degrees of freedom.

The 3D presentation of the positions and movement paths of occlusal or juxta-articular measurement points provides important information on the movement behavior of the mandibular joint and the teeth on the lower and/or upper jaw. The 3D presentation of distinctive positions in the face provides a mapping of the face symmetry for the fitting of the denture. In a functional, preliminary examination, any discoordinations and limitations of movement can be analyzed and documented.

The electronic position analysis of the condyles enables different occlusion positions to be compared and can thereby indicate possible pain vectors in the joint.

An analysis of the chewing movements takes place with a special software module. The XML export function enables the use of jaw movements determined with the SICAT JMT⁺ system in CAD/CAM systems and PACS systems for the functional optimization of dentures and of mouth-guards.

The measurement system may only be used by trained personnel, meaning dentists in dental facilities.

The SICAT JMT⁺ system is used to support the functional diagnosis. The measurement sensor technology consists of a receiving sensor and a transmitting sensor. This is attached to the patient's head. The lower jaw sensor technology is fitted with a special locking mechanism for fixing it to the attachment. The face bow is positioned at the front with the support on the nasion and at the back of the head above the ears. A measurement can then be carried out according to the required settings and measurement parameters in the software.

All the measurement and/or analysis results of the SICAT JMT⁺ system should always be interpreted by a trained specialist and checked for their relevance in consideration of the clinical medical history of the patient and in the context of the further diagnostic procedure. If invasive measures are taken, the measurement system should only be used as an additional assessment method. Under no circumstances can or should invasive surgery or measures that put the patient at risk be carried out based on the measurement results alone.

2.1.2 SICAT DATA EXPORT

The JMT export function enables the use of jaw movements determined with the SICAT JMT⁺ system in CAD/CAM systems and PACS systems for the functional optimization of dentures and of mouth-guards. The **SICAT Fusion Bite** serves as an interface for the referencing. It carries reference marks that can be captured by imaging systems such as surface scanners or DVT.

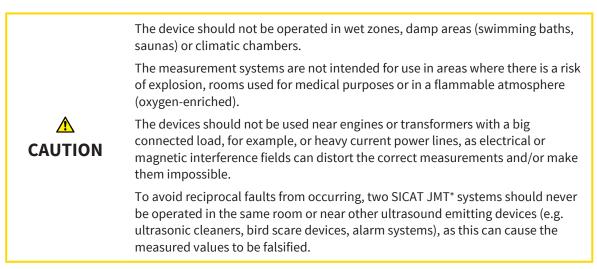
2.2 SAFETY

2.2.1 ENVIRONMENTAL CONDITIONS

The SICAT JMT⁺ jaw motion analysis system is suitable for use in dry interior rooms, as can be found in clinics, medical practices and laboratories.

Permissible operating temperature: 10°C to 40°C

Relative humidity: 30 % to 70 %



2.2.2 STORAGE AND TRANSPORT

The storage and transport of the measurement system should only be carried out in the original packaging, as provided by SICAT.

Storage temperature: -20°C to +70°C

Relative humidity: 5% to 90%

Air pressure: 700 hPa to 1060 hPa

2.2.3 OBLIGATIONS OF THE USER



The general guidelines and/or national legislation, national regulations and technical regulations pertaining to medical products are to be applied and fulfilled both with the start-up and during the operation of the SICAT product appropriate to the stated purpose. In Germany, operators, those responsible for such devices, and users are obliged to operate their devices in compliance with the MPG (Medical Devices Act) regulations.

- · It is the obligation of the user:
 - to comply with all the safety instructions stated in the instructions for use.
 - to carry out all of the inspection and maintenance work regularly as specified in the instructions for use.
 - to only use fault free working equipment.
 - to ensure that the device is functionally safe and in a proper state prior to every instance of use of the device.
 - to ensure all the provided instructions for use that form part of the measurement system, are accessible to all users at all time, and to keep them near the measurement system.
 - to protect oneself, the patients and third parties against dangers.
 - to prevent a contamination occurring due to the product.
- · During use, it is necessary to comply with the legal regulations, especially:
 - the current work safety regulations.
 - the current accident prevention measures.
- Responsibility is assumed to ensure the safety, reliability and effective performance of all measurement systems and accessories delivered by SICAT, such that:
 - assembly work, extensions, new settings, changes or repairs are carried out by SICAT or third parties authorized by SICAT, trained technicians or by the personnel of authorized dealers. The storage and transport should only be carried out in the original packaging, as provided by the manufacturer.
 - the product is operated in compliance with the instructions for use.
 - the information technology components provided by the operator comply with the technical requirements for hardware and software contained in these instructions for use, and that they are installed and set up according to the applicable descriptions for these components.
 - the place of installation corresponds with the specified environmental conditions for the measurement system and the current installation regulations.
 - only the software made available by SICAT, as well as the components and accessory parts listed in these instructions for use are used with the system.

2.2.4 GENERAL SAFETY INFORMATION



- The use and operation of the system and the evaluation of measurement data and its interpretation should only be carried out by trained specialist personnel. The manufacturer assumes no liability for damage to persons or property, or the loss of data that may occur due to the improper use of the software, the device, or its accessory parts.
- Patients and measurement data may only be copied, moved or deleted with the help of the database function that is provided by the SICAT application program. In the case of the deliberate changing of data without the database functions, the user alone bears the full risk.
- All measurement and/or analysis results should always be interpreted by a trained specialist and checked for their relevance in consideration of the clinical medical history of the patient and in the context of the further diagnostic procedure. If invasive measures are taken, the measurement system

should only be used as an additional assessment method. Under no circumstances can or should invasive surgery or measures that put the patient at risk be carried out based on the measurement results alone.

- In the case of malfunctions and/or defects being suspected and/or ascertained, the device has to be taken out of use immediately, labelled as 'Out of Use', and secured to prevent use. Please contact the manufacturer or your sales partner immediately under all circumstances.
- The measurement system has to be checked to ensure the measurement function is operating correctly at regular intervals. For further information about this, please see the *Functional checks*, *preparation*, *disposal* [> *Page 24*] chapter in these instructions for use.
- Do not install the jaw motion analysis system near a source of heat or in direct sunlight behind a window, as excessive heating can lead to incorrect measurement results.
- Ensure that all power and connection cables are routed and protected so that nobody is able to trip over them. Check all cables and connection plugs regularly for damage. Damaged power supply units, plug connectors and cables must be replaced before further use.
- The measurement system is not protected against the penetration of fluids. If fluid penetrates the measurement system, switch it off and please contact the SICAT GmbH & Co. KG technical service team.
- · Never introduce objects into components of the measurement system.
- Before starting every measurement, it is necessary to ensure the correct choice and correct position
 of the transmitters or application aids. The cables or the application aids can present a risk of injury
 to the patient. In this context, please consult the special instructions in the handbooks of the
 application software, and do not allow children or mentally impaired patients to enter the proximity
 of the device without supervision.

2.2.5 SAFETY INFORMATION ON HEART PACEMAKERS / DEFIBRILLATORS



- In the magnetic coupling for the attachment of the Lower jaw sensor type 24T on the Para-occlusal attachment 90 there are strong permanent magnets, such as those that are used on headphones on MP3 players. Under especially unfavorable circumstances, at short distances (< 15 cm), these magnets can have a negative impact on the functionality of certain implanted heart pacemakers and defibrillators. Therefore, the Lower jaw sensor type 24T should not be positioned on the upper body of the patient on patients with electronic implants.
- Version BT devices contain a Bluetooth transmitter as an interface to the PC. Although there is so far no evidence of a possible interference of heart pacemakers/defibrillators by Bluetooth transmitters, the SICAT JMT⁺ system is not recommended to be used on patients with electronic implants using the lanyard, but with the maintaining of a safety distance of at least 15 cm from the patient's thorax.
- No interference of electronic implants is to be expected from the ultrasonic transmitters used in the measurement system, as the SICAT JMT⁺ system works with airborne sound and a very low sound power of a few milliwatts. Due to the adverse connection during the transition from the air into the human body, the noise intensity of the measurement signals is weakened strongly such that any interference with implants, as well as any damage to tissue, is excluded.

2.2.6 PROHIBITED USE



· Improper and/or prohibited use of the measurement system is not permitted and express warning is herewith provided of such.

• Do not under any circumstances attempt to maintain or prepare the measurement system in any way other than as described in the instructions for use. This could cause the high sensitivity sensor technology to be in impaired terms of its measurement accuracy.

- In the case of malfunctions and/or defects being suspected and/or ascertained, the device has to be taken out of use immediately, labelled as 'Out of Use', and secured to prevent use, with the on/off switch being covered and secured with adhesive tape.
- Changing or modifying the measurement system or its accessory parts without the written permission of SICAT is not allowed. If the device is changed without permission, the operator is obliged to carry out suitable examinations and inspections in order to guarantee the secure use.
- SICAT measurement systems must not be operated in environmental conditions other than those stated in the *Technical data of the SICAT JMT***measurement system* [> *Page 11*] chapter (e.g. in an oxygen enriched environment, wet zones, damp rooms, climactic chambers, low pressure-, high pressure-, or altitude chambers, etc.).

3 PRODUCT DESCRIPTION

3.1 SYSTEM COMPONENTS

In the basic configuration, the SICAT JMT⁺ system consists of the following components:

- SICAT JMT⁺ basic unit
- · Lower jaw sensor type 24T (transmitter)
- Facebow type 13R (receiver)
- · USB power supply for supplying the measurement system for BT devices
- USB adapter for SICAT JMT⁺ (type A to bayonet connector, 3 m in length)
- · SICAT JMT⁺ application software

3.2 TECHNICAL DATA OF THE SICAT JMT⁺ MEASUREMENT SYSTEM

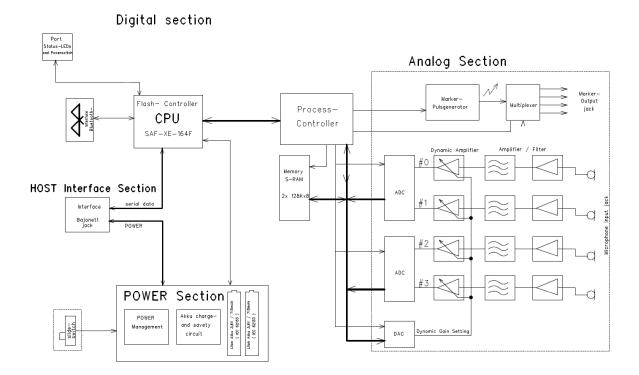
VERSION	SICAT JMT⁺ BT	
Dimensions (W x H x D)	111 x 86 x 31 mm	
Weight	205 g	
Power supply	5V DC / 1W (USB to charge battery)	
Battery	yes	
Measurement range	60 - 120 mm	
Ultrasonic frequency	40 kHz	
Max. measurement rate	50 Hz	
Positioning accuracy in the occlusal area	\pm 0.1 mm (y); \pm 0.2 mm (x, z)	
Interface	USB	
Bluetooth	yes	

3.3 MEASUREMENT PRINCIPLE OF THE SICAT JMT⁺ JAW MOTION ANALYSIS SYSTEM

The jaw motion analysis system consists of the **Lower jaw sensor type 24T** and the ultrasonic receiver module on the **Facebow type 13R**. The sensor technology components of the receiver and transmitter modules are each attached in geometrically defined positions. The marking points consist of small, sequentially operated ultrasonic transmitters. The accompanying facebow consists of six ultrasonic microphones. Both modules are connected with the evaluation electronics in the **SICAT JMT⁺ basic unit** via a connecting cable.

During operation, the ultrasonic transmitters continuously emit impulses, from the operating time of that, between the transmitter and receiver microphones, the evaluation electronic calculates the absolute room coordinates of the markers using a triangulation method.

The calculation of the measurement coordinates and further measurement parameters, as well as the compensation of the disturbance variables, occurs on a PC-supported basis in the evaluation programs.



BLOCK DIAGRAM OF THE MEASUREMENT SYSTEM

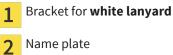
3.4 CONTROL ELEMENTS AND CONNECTIONS

VIEW FROM FRONT



VIEW FROM BELOW





3.5 MEANING OF THE DISPLAY LIGHTS

LED	ON/OFF SWITCH	MEANING			
Green / operational status indicator					
off	0 (off)	The measurement system is NOT in operation			
lit up	l (on)	The measurement system is in operation			
Orange / measurement					
off	l (on)	The measurement system is ini- tialized and is ready for meas- urement.			
flashes	l (on)	The measurement system is waiting for initialization, meas- urement not possible yet.			
lit up	l (on)	The measurement has started / ultrasonic transmitters are act- ive.			
Yellow / battery charging statu	5				
off	l (on)	USB adapter for SICAT JMT ⁺ and/or USB power supply con- nected, reduced charging as soon as charging status > 95%			
flashes	0 (off)	USB adapter for SICAT JMT ⁺ and/or USB power supply con- nected, battery charging, char- ging status < 95%			
	l (on)	Battery level critical < 20% Connect the USB adapter for SICAT JMT* or USB power sup- ply immediately, as it is pos- sible that data will be lost if the measurement is continued.			
lit up	0 (off)	USB adapter for SICAT JMT ⁺ and/or USB power supply con- nected, battery fully charged, charging status 100%			
Blue / Bluetooth connection					

LED	ON/OFF SWITCH	MEANING
off	l (on)	The measurement system is ini- tialized and is ready for meas- urement.
lit up	l (on)	The measurement has started / the measurement system is connected with the PC via Bluetooth

3.6 ASSIGNMENT OF THE CONNECTING SOCKETS

LOWER JAW SENSOR TYPE 24T / DIGITAL INPUT (LIGHT GREEN)

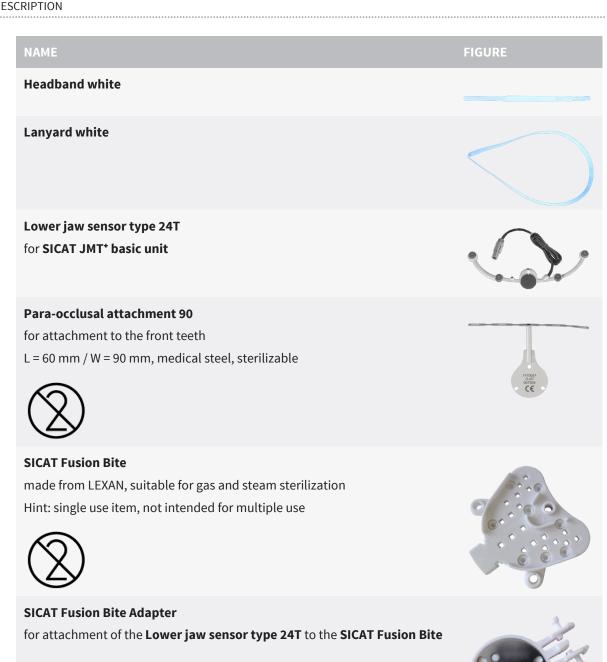
SIGNAL	PIN
Marker 1	Pin 1
Marker 2	Pin 2
Marker 3	Pin 3
Marker 4	Pin 4
SSWBus	Pin 5
n.c.	Pin 6
SDA (I ² C)	Pin 7
Signal	Pin
SCL (I ² C)	Pin 8
+3.3 Volt	Pin 9
DRY_A	Pin 10
DRY_G	Pin 11
Dig. Input	Pin 12
GND	Pin 13
GND	Pin 14

FACEBOW TYPE 13R / DIGITAL INPUT (WHITE)

SIGNAL	PIN
Microphone 1	Pin 1
Microphone 2	Pin 2
Microphone 3	Pin 3
Microphone 4	Pin 4
SSWBus	Pin 5
Mic-select	Pin 6
SDA (I ² C)	Pin 7
SCL (I ² C)	Pin 8
+3.6 - 12 Volt	Pin 9
DRY_A	Pin 10
DRY_G	Pin 11
Dig. Input	Pin 12
GND	Pin 13
GND	Pin 14

3.7 ACCESSORIES AND SPARE PARTS





USB power supply for charging the SICAT JMT⁺ basic unit



NAME

EU - Adapter **USB power supply** UK - Adapter **USB power supply** USA - Adapter **USB power supply** Australia - Adapter **USB power supply** World - Adapter **USB power supply**

USB adapter for SICAT JMT⁺ Data connection of measurement system and PC



SICAT JMT⁺ USB card

Medium that contains the SICAT JMT⁺ software



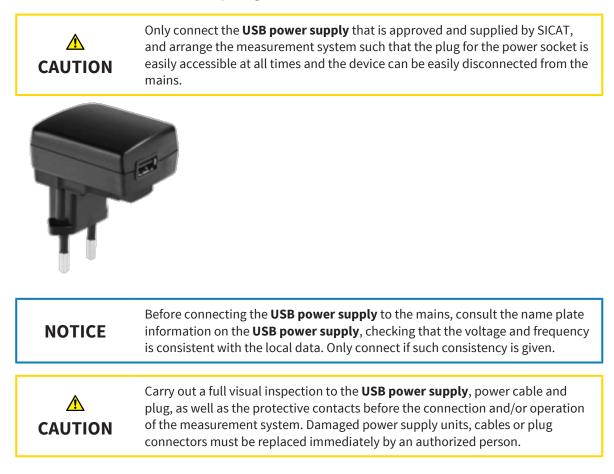
4 PUTTING THE MEASUREMENT SYSTEM INTO OPERATION

For the commissioning of the jaw motion analysis system, a USB cable, as well as the medium with the SICAT JMT⁺ application software are required. All components are included in the scope of delivery for the SICAT JMT⁺ system.

4.1 POWER SUPPLY AND CHARGING THE BATTERY

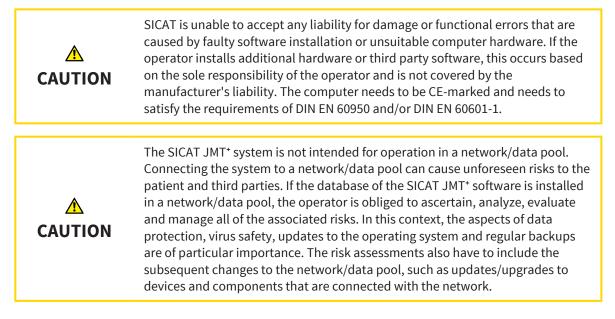
For the rapid charging of the battery of the **SICAT JMT⁺ basic unit** when it is in the deactivated state, connect the **USB power supply** with an AC outlet and a USB cable with the USB socket on the **SICAT JMT⁺ basic unit**.

Alternatively, the **SICAT JMT⁺ basic unit** can also be charged or operated directly on the USB socket of a PC. To do this, connect the PC directly using a USB cable.



4.2 COMPUTER REQUIREMENTS

For the requirements of the SICAT JMT⁺ system with concern to a PC/laptop, please refer to the instructions for use on SICAT JMT⁺ software.



4.3 INSTALLING THE SICAT JMT⁺ SOFTWARE

Installation instructions are provided in the instructions for use on the SICAT JMT⁺ software.

NOTICE Please make sure under all circumstances, that before connecting the measurement system and PC with the USB cable and/or via Bluetooth that you have installed the SICAT JMT⁺ software.

If this step is omitted, problems can occur during the installation of the device driver, as the Windows operating system registers the location of the driver on the hard disk on the occasion of the first connection of the SICAT JMT⁺ system and the PC.

If no corresponding SICAT JMT⁺ software is installed on the PC yet, the mapping of the driver will fail for the above reasons, and the SICAT JMT⁺ system may not function correctly.

NOTICE If problems should occur with the hardware driver of the SICAT JMT⁺ system, please unplug the USB cable from the PC and start it again. Now, re-install the SICAT JMT⁺ software and create the connection once again.

4.4 CONNECTING THE ACCESSORY PARTS

Connect the variable and variable with the corresponding colored sockets on the **Facebow type 13R**. (Images see *Control elements and connections* [> *Page 13*])

NOTICE When connecting the variable and variable to the **Facebow type 13R**, please ensure that the plugs are protected against false polarity / being plugged into the wrong socket with a keying nose. All of the plugs should go into the sockets easily and without much effort.

Next, connect the **SICAT JMT⁺ basic unit** and a USB interface on your computer with the USB cable provided, or set up the Bluetooth connection. In this context, it is necessary to ensure that the measurement system is switched on. Your measurement system is now ready to use. Detailed instructions on the operation of the SICAT JMT⁺ system are provided in the instructions for use for the SICAT JMT⁺ software.

4.5 TAKING THE MEASUREMENT SYSTEM OUT OF OPERATION

To take the measurement system out of operation, please start by connecting the SICAT JMT⁺ software, shutting the PC down, and switching it off. Next, switch the **SICAT JMT⁺ basic unit** off and decouple the USB connection from the PC or **USB power supply**. Next, unplug the **USB power supply** from the socket.

5 FUNCTIONAL CHECKS, PREPARATION, DISPOSAL



 Regular maintenance and care of the measurement system helps to prevent damage and guarantees its long-term safety. All of the procedures stated in these instructions for use regarding the maintenance and preparation of the system are to be carried out on a regular basis.

- If the measurement system or accessory parts show damage, they should be sent to the manufacturer for a safety inspection. In the case of malfunctions and/or defects being suspected and/or ascertained, the device has to be taken out of use immediately, labelled as 'Out of Use'.
- All maintenance and servicing work on the measurement system or individual parts that are extended beyond the activities described in the instructions for use may only be completed by SICAT or persons authorized by SICAT.
- Before beginning the preparation, switch the measurement system off under all circumstances and disconnect it completely from the power supply network.

5.1 SPECIFIED REGULAR CHECKS AND TECHNICAL SAFETY CHECKS



- SICAT GmbH & Co. KG does not specify any technical safety checks for the SICAT JMT⁺ system.
- To maintain the proper status of the electrical operating equipment, it is necessary to carry out repeated inspections and technical safety checks (e.g. in Germany according to BGV A3 accident prevention regulations as well as technical safety checks pursuant to the German Medical Products Operator Ordinance). In this context, it is necessary to ensure that these are not SICAT -specific measures, but the current regulations regarding electrical devices.
- For safety reasons, inspecting the proper state of all the connection cables, network cables, USB
 power supply and power sockets is recommended before every use of the measurement system. If
 parts are damaged it is necessary to replace them before using the measurement system again.
- · Immediate servicing measures are to be carried out by SICAT if:
 - any liquids/fluids entered the device.
 - the cables or plug connectors show any damage.
 - parts of the sensor technology are damaged.
 - covers are damaged or have come off.
 - a defect or fault is either suspected or ascertained.
- · If the name plate or other labelling (e.g. warning notices) on the machine are damaged or unreadable, these are to be replaced.

CAUTION

5.2 CHECKING THE MEASUREMENT FUNCTION

To guarantee long-term patient safety, the SICAT JMT⁺ system has to be inspected at regular intervals to ensure its proper measurement function.

After hard knocks, such as if the **Facebow type 13R** or **Lower jaw sensor type 24T** should fall on the floor, it is necessary to check the measurement function immediately.

In the event of evident damage to system components (warping, dents, cracks), no further measurements should be carried out.

- The ultrasonic transmitters of the **Lower jaw sensor type 24T** can be inspected for their function during a measurement by listening to see whether a regular cracking is emitted from each of the transmitters.
- To inspect the system, for known jaw functioning measurements (e.g. known maximum opening width, known condylar range of motion with protrusion, known horizontal condylar guidance inclination), the user can measure himself with the measurement system. These measurement results should correspond with the known values.
- When the sensors do not move, the SICAT JMT⁺ software should show an unmoving image of the lower jaw. Possible deviations (spikes or jumps in the measurement curve in spite of unmoved markers, incorrect presentation of the lower jaw, etc.) indicate a faulty measurement and impair the evaluation.
- Should doubts surround the measurement accuracy, inspecting the SICAT JMT⁺ system at SICAT is recommended in order to ensure the stated measurement accuracy.

5.3 RECTIFYING FAULTS

In the event of faults, please start by checking the following points:

- Is the SICAT JMT⁺ system switched on and being supplied with electricity? (green operating display LED is lit on the measurement system, batteries are charged or **USB power supply** and/or USB cable is connected)
- Has the USB connection and/or Bluetooth connection between the measurement system and measuring PC been made correctly?
- Are all the other components in the measurement system (Facebow type 13R, Lower jaw sensor type 24T) connected correctly?

NOTICE For further information on error messages and their rectification, please refer to the instructions for use on SICAT JMT⁺ software.

CHECK LIST FOR THE RECORDING OF ERROR MESSAGES

NOTICE

To be able to provide you with the optimum support in the case of operational faults to your SICAT JMT⁺ system, our service team employees require the following information:

- Serial numbers of the SICAT JMT⁺ basic unit and Lower jaw sensor type 24T/Facebow type 13R
 - The serial numbers are on the name plates on the rear of the SICAT JMT⁺ basic unit and/or on the cables for the Facebow type 13R and Lower jaw sensor type 24T.
- · SICAT JMT⁺ software version
- Operating system version of your measurement PC
 - e.g. Windows 7 Professional Service pack 1 (to find under Windows 7: Windows Start button > Control Panel > System)
- · Further components connected to the measurement system
- · List of all USB/Bluetooth devices connected to the measurement PC
 - e.g. mouse, printer, other measuring systems etc.
- \cdot $\,$ Screen shot of the error message, or exact wording
 - e.g. "Timeout reading from USB"
- · Accurate as possible description of the process that led to the error message
 - e.g. "Type A" measurement started, button "B" then clicked, movement "C" then completed, changed over to function "D", upon changing back, the error message "xyz" was reported, etc.

5.4 PREPARATION METHODS

NOTICE After every case of use of the SICAT JMT* system, a re-preparation is required according to DIN EN ISO 17664. All accessory parts that come into contact with the patient's mucosa have to be sterilized before use. Before starting cleaning work or disinfection, switch the measurement system

Before starting cleaning work or disinfection, switch the measurement syster off under all circumstances and disconnect it completely from the USB connection and the USB power supply.



The following accessory parts are only intended for one-off use on a patient, and should not be prepared again after use.

- · Para-occlusal attachment 90
- · SICAT Fusion Bite

5.4.1 MANUAL CLEANING

- Before sterilization, clean the accessory parts by hand under running water (drinking water quality, $30 \degree C \pm 5 \degree C$, flow rate 2 liters/min.) with a medium strength toothbrush for 30 seconds.
- · Complete the sterilization immediately subsequent to the cleaning.
- The cleaning of measurement systems and electrical accessories (**Facebow type 13R**, **Lower jaw sensor type 24T**) should only be carried out when the system is switched off and the **USB power supply** and/or USB cable are unplugged, and using a damp cloth.

5.4.2 MANUAL DISINFECTION

The electrical system components can be wipe-disinfected with suitable solutions. Disinfect all electric components (**Facebow type 13R**, **Lower jaw sensor type 24T**) with a cloth that has been dampened with a disinfectant solution.



RECOMMENDED DISINFECTANT SOLUTION

Composition approx. 25% ethanol, 35% propanol

e.g. "Mikrozid "Liquid / Schülke & Mayr; Original CaviCide / Kerr Corporation or comparable solutions

NOTICE	When using a disinfectant solution, please comply with the recommendations stated by the manufacturer, especially the specified application time.
▲ WARNING	Due to the risk of possible mix-ups, chemicals that are required for the disinfection or cleaning must only be stored, prepared and kept ready in their intended containers.

5.4.3 STERILIZATION

All accessory parts that come into contact with the patient's mucosa have to be sterilized before use.

NOTICE The sterilization is to be completed immediately subsequent to the cleaning.

Sterilize the bite-forks and lower jaw attachment with a fractionated pre-vacuum for four minutes at 134 °C \pm 1 °C and 2 bar (can be sterilized up to a max. of 138 °C).

It is necessary to sterilize the following accessory parts:

- · Para-occlusal attachment 90
- SICAT Fusion Bite

5.5 DISPOSAL

5.5.1 PACKAGING

In Germany, all of the transport packaging supplied by SICAT can be disposed of into the materials cycle at local waste disposal centers.

5.5.2 DISPOSAL OF ELECTRONIC WASTE



The adjacent symbol shows that pursuant to the EC Directive on Waste Electrical and Electronic Devices (2012/19/EU) and national legislation, a product cannot be disposed of via the household waste, and must be disposed of separately within Europe.

For this purpose, at the end of its useful life, the measurement system can be returned by the customer to SICAT GmbH & Co. KG, and will then be forwarded by SICAT at no extra cost and without reimbursement to the appropriate recycling companies.

Due to potentially dangerous substances that can often be found in electrical or electronic devices, the improper handling of old devices can have a negative impact on the environment and human health. In disposing of this product properly, you also contribute to an effective use of natural resources.

5.5.3 ACCUMULATORS AND BATTERIES

Accumulators and batteries do not belong in the household waste! In the interests of protecting the environment, the end consumer is legally obliged (Battery Directive) to hand in old and used batteries. Used batteries and power packs can be handed in at the district collection points, or in all locations where such batteries are offered for sale. The batteries are accepted at no cost to the consumer.

6 SAFETY STANDARDS AND CLASSIFICATION OF THE SYSTEM

6.1 CLASSIFICATION PURSUANT TO APPENDIX IX OF THE DIRECTIVE 93/42/EEC

The system is classified as a medical product of **Class I with a measuring function**.

6.2 SAFETY OF ELECTRICAL MEDICAL DEVICES

The device fulfils the requirements of the DIN EN 60601-1:2006 standard.

Classification corresponding to DIN EN 60601-1

Type BF

Protection class II

Continuous operation

Not suitable for use in an oxygen-enriched environment

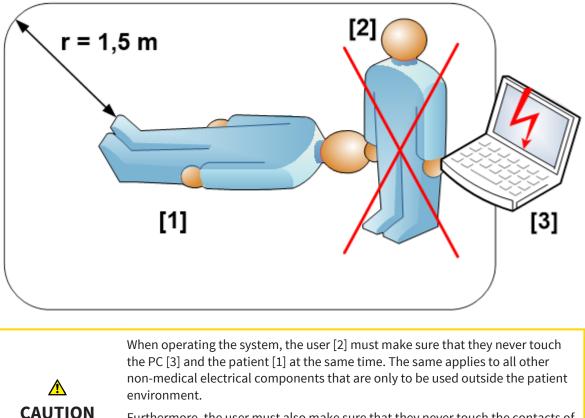
6.2.1 CONNECTION OF THE SICAT JMT⁺ MEASUREMENT SYSTEM WITH OTHER ELECTRICAL DEVICES

(also see DIN EN 60601-1:2006, section 16 Medical Electrical Systems)

▲ CAUTION	The SICAT JMT ⁺ system may only be connected with other electrical devices if they correspond with the requirements of DIN EN 60950 and/or DIN EN 60601-1 or have been designated as compatible by SICAT GmbH & Co. KG.		
	When connecting several devices to a single measurement station, it is necessary to ensure that no dangers can occur due to the accumulation of leakage currents.		
A CAUTION	Devices with that the patient comes directly into contact, and that are used together in a single medical electrical system, must comply in complete form with all the requirements of DIN EN 60601-1:2006, section 11.		
	There is the risk of an electrical shock upon contact with devices that are not earthed separately.		

6.2.2 ENVIRONMENT OF THE PATIENT / TEST PERSON

In practice, an experience value with a clearance of 1.5 m from the patient has proven effective for the determination of the patient environment.



Furthermore, the user must also make sure that they never touch the contacts of the measurement system plug connectors and the patient at the same time.

Failure to comply can lead to the occurrence of dangerous leakage currents.

The following components of the SICAT JMT⁺ system may only be used within the patient environment:

· SICAT JMT⁺ basic unit including sensor technology



The computer and other non-medical electrical accessories have to be set up outside of the patient environment (1.5 m clearance).

6.3 ELECTROMAGNETIC COMPATIBILITY FOR GUIDELINES / MANUFACTURER DECLARATION

The SICAT JMT⁺ system satisfies the requirements of the EN 60601-1-2 standard. (Medical electrical devices - part 1-2: General concerning for safety including the key performance attributes - supplementary norm: Electromagnetic compatibility - requirements and tests).

Testing authority:

Schwille Elektronik in Munich

Produktions- und Vertriebs GmbH

Benzstrasse 1A

85551 Kirchheim

Detailed information on EMC values and manufacturer data is provided in the tables for this chapter of the instructions for use.

Medical electrical devices are subject to special safety measures regarding EMC (electromagnetic compatibility) and have to be installed and put into operation pursuant to the instructions provided below.

▲ CAUTION	Although the SICAT JMT ⁺ system corresponds with the requirements of the DIN EN 60601-1-2 norm in terms of all points, mobile phones cannot be excluded from influencing the jaw motion analysis system. If possible, such devices should not be operated during measurements in the environment of the JMT measurement system.
▲ CAUTION	The use of accessories, especially connection cables to the PC that are not supplied by SICAT for the SICAT JMT ⁺ system, or are expressly approved for use with the device, can lead the JMT systems to have increased emissions or reduced interference immunity.
▲ CAUTION	The SICAT JMT ⁺ system should not be used near X-ray systems, engines or transformers with a big connected load, for example, as electrical or magnetic interference fields can distort the correct measurements and/or make them impossible. The same applies to neighboring power cables and devices that do not have CE marking. If operation is required in the immediate proximity of sources of interference, it is necessary to observe the device under all circumstances to make sure it is operating in the proper way in this arrangement.

GUIDELINES AND MANUFACTURER'S DECLARATION - ELECTROMAGNETIC EMISSIONS

The measurement systems in the JMT range of products are intended for operation in the electromagnetic environment stated below. The customer or user of the JMT measurement system should ensure that it is operated in such an environment.

Emissions tests	Compliance	Electromagnetic environment - guideline
RF emissions according to CISPR 11	Group 1	The measurement systems in the JMT range of products only use RF energy for their internal functioning. For this reason, their RF emissions are very low and it is unlikely that neighboring electronic devices will suffer interference.
RF emissions according to CISPR 11	Class B	The JMT measurement system is suitable for u in all facilities, including those in domestic set
Harmonic emissions according to IEC 61000-3-2	Class B	tings and those directly connected to the public power supply network that supplies buildings that are used for domestic purposes.
Voltage fluctuations/ flicker according to IEC61000-3-3	Compliance	

GUIDELINES AND MANUFACTURER'S DECLARATION - ELECTROMAGNETIC INTERFERENCE IMMUNITY

The measurement systems in the JMT range of products are intended for operation in the electromagnetic environment stated below. The customer or user of the JMT measurement system should ensure that it is used in such an environment.

Immunity tests	IEC 60601- test level	Compliance level	Electromagnetic environment - guidelines
Electrostatic dis- charge (ESD) ac- cording to IEC 61000-4-2	± 6 kV contact dis- charge ± 8 kV air dis- charge	± 6 kV contact dis- charge ± 8 kV air dis- charge	Floors should be made from wood or concrete or fitted with ceramic tiles. If the floor is equipped with synthetic material, the relative air humidity should be at least 30%.
electrical fast transient/bursts according to IEC 61000-4-4	± 2 kV for power cables ± 1 kV for input and output cables	± 2 kV for power cables ± 1 kV for input and output cables	The supply voltage quality should be that of a typical commercial or hospital environment.
Surges according to IEC 61000-4-5	± 1 kV differential mode ± 2 kV common mode	± 1 kV differential mode ± 2 kV common mode	The supply voltage quality should be that of a typical commercial or hospital environment.

GUIDELINES AND MANUFACTURER'S DECLARATION - ELECTROMAGNETIC INTERFERENCE IMMUNITY

Voltage dips, short interrup- tions and voltage variations on the supply voltage ac- cording to IEC 61000-4-11	< 5% U_{T} (> 95% drop to the U_{T}) for ½ a period 40% U_{T} (60% drop to the U_{T}) for 5 periods 70% U_{T} (30% drop to the U_{T}) for 25 periods < 5% U_{T} (> 95% drop to the U_{T}) for 5 s	< 5% U_{τ} (> 95% drop to the U_{τ}) for $\frac{1}{2}$ a period 40% U_{τ} (60% drop to the U_{τ}) for 5 periods 70% U_{τ} (30% drop to the U_{τ}) for 25 periods < 5% U_{τ} (> 95% drop to the U_{τ}) for 5 s	The supply voltage quality should be that of a typical commercial or hospital environment. If the user of the meas- urement systems in the JMT range of products requires continued function- ing during power supply interruptions, it is recommended that the JMT jaw motion analysis system be powered by an interruption-free power supply or a battery.
Magnetic field with the power supply frequency (50/60 Hz) accord- ing to IEC 61000-4-8	3 A/m	3 A/m	Power frequency magnetic fields should correspond with the typical val- ues that can be found in the commer- cial or hospital environment.

NOTE U_T is the A.C. mains voltage prior to application of the test level.

GUIDELINES AND MANUFACTURER'S DECLARATION - ELECTROMAGNETIC INTERFERENCE IMMUNITY

The measurement systems in the JMT range of products are intended for operation in the electromagnetic environment stated below. The customer or user of the JMT measurement system should ensure that it is used in such an environment.

Immunity tests	IEC 60601- test level	Compliance level	Electromagnetic environment - guidelines
			Portable and mobile radio equipment should be used no closer to the JMT measurement system, including the cables, than the recommended safety distance, calculated from the equation applicable to the frequency of the transmitter. Recommended safety distance:
Conducted RF surges according to IEC 61000-4-6	3 V _{eff} 150 kHz to 80 MHz	3 V _{eff}	$d = 1, 2\sqrt{P}$
Radiated RF surges according to IEC 61000-4-3	3 V/m 80 MHz to 2.5 GHz	3 V/m	$d = 1, 2\sqrt{P}$ for 80 MHz to 800 MHz $d = 2, 3\sqrt{P}$ for 800 MHz to 2.5 GHz

GUIDELINES AND MANUFACTURER'S DECLARATION - ELECTROMAGNETIC INTERFERENCE IMMUNITY

Where P is the rated output of the transmitter in watts (W) according to the information of the transmitter manufacturer and d is the recommended safety distance in meters (m).

The field strength from fixed RF transmitters as determined by an electromagnetic site survey ^a is less than the compliance level in all frequency ranges. ^b

Interference may be possible in the vicinity of equipment that is marked with the following symbol



NOTE 1 At 80 MHz and 800 MHz, the higher value applies.

NOTE 2 These guidelines may not apply in all cases. Electromagnetic propagation is affected by absorption and reflection from buildings, objects and people.

a The field strengths of fixed transmitters, such as base stations for radio telephones and land mobile radios, amateur radio, AM and FM radio broadcasters and TV broadcasters, cannot be theoretically predicted with accuracy. To assess the electromagnetic environment due to the fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength at the location of the JMT measurement system exceeds the aforementioned compliance level, the JMT measurement system should be observed to verify its proper functioning. If abnormal performance attributes are observed, additional measures can be necessary, such as re-orientation or an alternative location of the measurement system of the JMT range of products.

b Over the frequency range of 150 kHz to 80 MHz, the frequency range should be lower than 3 V/m.

RECOMMENDED SAFETY DISTANCES BETWEEN PORTABLE AND MOBILE RF COMMUNICATION DEVICES AND THE MEASUREMENT SYSTEMS IN THE JMT RANGE OF PRODUCTS

The measurement systems in the JMT range of products are intended for operation in an electromagnetic environment in that radiated RF interference is controlled. The customer or the user of the JMT measurement system can help prevent electromagnetic interference by maintaining a minimum clearance between portable and mobile RF communication devices (transmitters) and the JMT measurement system - depending on the output power of the communication device as stated below.

Rated output of trans- mitter W	Safety distance depending on the transmitter frequency m			
	150 kHz to 80 MHz	80 MHz to 800 MHz	800 MHz to 2.5 GHz	
	$d = 1, 2\sqrt{P}$	$d = 1, 2\sqrt{P}$	$d = 2, 3\sqrt{P}$	
0.01	0.12	0.12	0.23	
0.1	0.38	0.38	0.73	
1	1.2	1.2	2.3	
10	3.8	3.8	7.3	
100	12	12	23	

For transmitters whose maximum rated output is not stated in the above table, the recommended safety distance d in meters (m) can be estimated using the equation applicable to the corresponding column, where P is the rated output of the transmitter in watts (W) according to the information of the transmitter manufacturer.

NOTE 1 At 80 MHz and 800 MHz, the higher frequency range applies.

NOTE 2 These guidelines may not apply in all cases. Electromagnetic propagation is affected by absorption and reflection from buildings, objects and people.

GLOSSARY

SICAT JMT⁺

SICAT JMT $\mbox{ records the motion of the lower jaw.}$

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