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CM LOC[®] Abutment

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Instructions for Use CM LOC® Abutment

1 Scope of application of Instructions for Use

These Instructions for Use apply to the products listed under Section 29. The issuing of these Instructions for Use renders all previous versions invalid. The manufacturer rejects any liability for damages resulting from non-compliance with these Instructions for Use.

2 Trade name

See Section 29.

3 Intended use

The products are intended for prosthetic restorations and to support procedures in the dental clinic or laboratory.

4 Expected clinical benefit

Restoration of chewing function and improved aesthetics.

The Summary of Safety and Clinical Performance, SSCP for the implantable devices covered by these Instructions for Use, is available on our website and accessible at this address: www.cmsa.ch/docs.

5 Product description

Product	Description
	CM LOC® Abutment Implant anchorage for hybrid-supported removable dental prostheses on implants, in combination with the specific CM LOC® system for female parts.
	CM LOC® Female part Retaining element as connecting part between denture and abutment.
 <p>extra-low ca. 400g low ca. 1200g medium ca. 1800g strong ca. 2400g</p>	CM LOC® Retention inserts Exchangeable retention inserts in four defined force levels. yellow: extra low red: low green: medium blue: strong
	CM LOC® CAD/CAM Retention element Additional retention element on CAD/CAM milled dental bars, in combination with the specific CM LOC® system for female parts.

6 Indications

CM LOC® Abutment:

Implant anchorage of hybrid-supported removable dental prostheses on implants, in combination with the specific CM LOC® system for female parts.

Mandible

CM LOC® Abutment:

Anchorage of mandibular (MD) prosthesis on 2 or more implants.

Maxilla

CM LOC® Abutment:

Anchorage of maxillary (MX) prosthesis on 4 or more implants.

CM LOC® CAD/CAM Retention element:

As additional retention element on CAD/CAM milled dental bars, in combination with the specific CM LOC® system for female parts.

7 Contraindications

- Implant divergences > 20° (per implant).
- The CM LOC® Abutments are to be used exclusively with the implant systems listed in Section 29.
- Use on a single implant.
- Not suitable if a fixed connection is required.
- Unilateral free-end prosthesis without transversal support.
- Use on root canal caps.
- Immediate restoration if immediate loading is not indicated for the implant.
- Implant system is not approved for the application.
- For additional contraindications, please refer to the instructions for use from the implant manufacturer.
- Patients who are unable to keep the regularly required check-up appointments for health reasons.
- Patients with bruxism or other para-functional habits.
- Patients with allergies to materials used in the product, see Section 19.
- Existing clinical picture in the patient's mouth does not permit the correct application of the products.

8 Compatible products

The CM LOC® Abutments are compatible with several implant interfaces and may only be combined with the compatible implant system.

The following CM LOC® specific components can be used for the application.

- System for female parts.
 - All auxiliary tools and auxiliary instruments.
- Exception screwdriver: the CM LOC® and CM LOC® FLEX Abutments have different, specific screwdrivers.

The CM LOC® system for female parts is occasionally compatible with the following Locator®-like abutments:

- CM LOC®
- CM LOC® FLEX
- MedentiLOC®

Please contact us for further information regarding other compatibilities.

To fabricate the finished denture, a number of general laboratory supplies are required in addition to the products listed under Section 29. The following gives a selection of materials that Cendres+Métaux SA offers in its portfolio.

08052138	Polyurock Kit	08055014	Livento® invest Powder (50 x 100 g)
08052135	Polyurock Catalyst	083739	Livento® invest Liquid (1000 ml)
08052136	Polyurock Release Spray	08052160	uniVest® Plus Powder (30 x 150 g)
08052137	Polyurock Mixer	08052161	uniVest® Plus Liquid (1000 ml)
08052566	Polyurock Colour yellow	08052162	uniVest® Rapid Powder (30 x 150 g)
08052149	ABF Wax Universal	08052163	uniVest® Rapid Liquid (1000 ml)
08052150	ABF Wax Creativ light	080181	CM soldering investment (4 kg)
08052151	ABF Wax Creativ dark	080229	CM soldering paste
08052154	ABF Wax Special	08052307	Legabril Diamond (50 g)
08052148	ABF Wax Margin		
08052153	ABF Wax Position		
08052152	ABF Wax Tecno		

9 Qualification of the specialist

Expertise in professional dentistry and dental technology is assumed. The current Instructions for Use must be available at all times and be completely read and understood before the first application. The fabrication of dentures and their maintenance may only be performed by qualified specialists.



Important information for the specialist



Warning symbol for increased caution

10 Prescription

Federal laws in the USA prohibit the use by or sale to unlicensed dentists.

11 Side effects

This product must not be used in patients with allergies or suspected allergies to materials used in the product (see Section 19), or only after prior allergological clarification.

Auxiliary instruments may contain nickel.

If applied as intended, side effects can be excluded.

12 Warnings**Magnetic resonance (MR) environment**

The device has not been evaluated for safety and compatibility in the MR environment.

The product has not been tested for heating or migration in the MR environment.

CM LOC® Spacer

The CM LOC® Spacer is slightly oversized with regard to the original components.

This ensures optimal space conditions for later polymerisation in the mouth.

The spacer must not be used in place of the female part or as a temporary replacement.

13 General information

-  – The height of the CM LOC® Block-out spacer can be shortened when using different abutment heights to achieve better mounting of the female part.
- When working with the CM LOC® Abutment for retention of overdentures, the direct or indirect approach may be used.
- We recommend that the clinical case be designed such that the largest possible support polygon is achieved. Small distances between consecutive implants and long free-end saddles can cause undesirable effects such as increased wear of the system components.
- Proper seating of the dentures on the mucosa must be checked at least once each year, and relining carried out if required to prevent rocking movement (overload). We recommend checking the prosthesis at regular intervals of approx. 3 months and to replace the retention inserts if necessary.
- When fabricating new dentures and when using a palate-free design, we recommend fabricating an individual reinforcement framework.
-  – In patients with suspected titanium allergy or hypersensitivity, we alternatively recommend the use of the Pekkton® female part. One must allow for an increased aftercare effort and, if necessary, changing/replacing the system for female parts, as Pekkton® is somewhat softer than titanium as the material for the female part.

Integration of the female part housing**Direct method**

The dentist providing treatment may integrate the CM LOC® housing of the female part and the retention inserts in an existing or new prosthesis directly during the treatment session.

Indirect method

The dentist must take an impression part of the CM LOC® Abutment and send the impression to the laboratory for subsequent fabrication of the model. The laboratory then inserts the CM LOC® Analog in the CM LOC® Impression part to facilitate reliable transfer of the position of the CM LOC® FLEX Abutment in the mouth, and fabricates the master model.

14 Preventive measures

-  – Screw in the product only once using the torque specified for this purpose.
- In case of immediate loading (observe implant manufacturer's indication) ensure that the tightening torque of the abutment does not exceed the torque of the implant > recommendation 5 Ncm below the tightening torque of the implant.
-  – The product components are supplied non-sterile. For more information see Section 16 "Reprocessing".
- Only original tools and parts may be used for this work. For information and additional details, please contact your Cendres+Métaux SA representative.
- Before any procedure, ensure that all required product components are available in sufficient quantity.
- For your own safety, always wear suitable protective clothing. In particular when grinding, we recommend wearing protective goggles and a dust mask as well as the use of a suction unit.
- Secure parts against aspiration.
- The mechanical cleaning by patients with a toothbrush and toothpaste may lead to premature wear.

15 Single use

Products that are intended for single use and are labelled "single-use" accordingly are subject to a certain amount of stress, increased wear, and even loss of functionality during their use.

-  Multiple application of products labelled «single use» was not tested. This can impair the safety, function and performance of the products as well as increase the risk of transmitting infections.

16 Reprocessing

-  The prosthetic work, including all system components, must be cleaned, disinfected and, if appropriate, sterilised prior to each work step. Materials made of metal alloys, high-performance polymers (Pekkton®) and ceramics are suitable for steam sterilisation. With the exception of Pekkton®, components made of plastics are not suitable for steam sterilisation. Consider published national guidelines when selecting a disinfection and sterilisation process and the Instructions for Use "Reprocessing of surgical and prosthetic products" (www.cmsa.ch/docs).

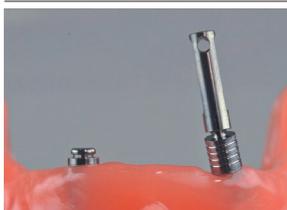
17 Scope of application

The CM LOC® Abutment components are designed to fixate overdentures (full dentures) or partial dentures completely or partially through endosseous implants with the specific CM LOC® System for female parts.

The CM LOC® Anchoring system consists of a standardised abutment with which implant divergences of up to 40° are indicated and a system for female parts with four exchangeable retention inserts in four defined force levels.

18 Procedure**Fabrication of a new prosthesis with CM LOC® Abutment.**

Patient Situation, Initial Position.

**Determining the implant axis**

Use the CM LOC® Case Guide to determine the divergence of the implant axes between the implants. Place the CM LOC® Case Guide on the implant for this purpose by screwing in manually.

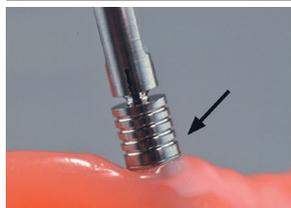
-  A corresponding CM LOC® Case Guide is available for each implant system.



By cyclically tipping until it stops (20°), use the CM LOC® Case Guide to determine the implant axes, so that the individual implant axes to each other can be determined.

 View from side and front.

 Should it not be possible to align the CM LOC® Case Guides parallel, a divergence of 20° between the implants is exceeded. If the divergence is greater than 20°, the CM LOC® Abutment may not be used, instead the CM LOC® FLEX is used.



Determining the abutment height

Choose the abutment height based on the implant position/gingival height and read off based on the graduation marks on the CM LOC® Case Guide. Determine the correct height of the CM LOC® Abutment with the lower edge of the CM LOC® Abutment positioned at least 1 mm above the gingiva. The lowest height starts at graduation marking 1.



Insert the CM LOC® Abutment

First, place the CM LOC® Abutment on the CM LOC® Screw Driver and screw it into the implant by hand.



Use the torque ratchet to tighten to the required torque. Make sure that the CM LOC® Screwdriver is correctly seated on the abutment. Secure all parts against aspiration.

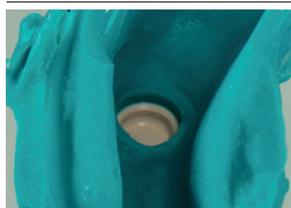
After assembly, the CM LOC® Screwdriver can be removed laterally by turning it back slightly.

 The screwdriver features an ISO connection and fits onto the coupling inserts for the corresponding torque wrenches.

Impression taking of the oral situation for further processing using the indirect method



Place the CM LOC® Impression part on the CM LOC® Abutment and create a functional impression. Ensure that the CM LOC® Impression part is correctly seated. Use a solid impression material (e.g. Impregum™).



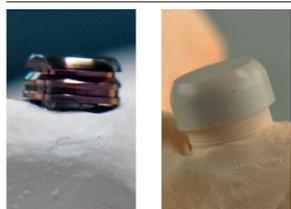
 Check that the material is fully distributed around the CM LOC® Impression part and that no impression material has spilled into the CM LOC® Impression part.

Otherwise, clean the abutment and repeat the impression-taking process.



Then pass to dental laboratory for fabrication of the model.

To fabricate the model in the laboratory, place the CM LOC® Analog in the CM LOC® Impression part and fabricate the master model.



Then place either the CM LOC® Housing of the female part with a mounted CM LOC® Processing insert or the CM LOC® Spacer onto the CM LOC® Analog.

Use of the CM LOC® Spacer or the original CM LOC® Housing of the female part is at the discretion of the user.

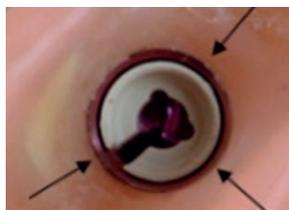
 Spacer = place holder for polymerisation in mouth.
Direct polymerisation with CM LOC® Housing of the female part in the laboratory.
Block out all undercuts during further processing and fabrication of the denture.



Finalisation

The prosthesis can now be fabricated using conventional technology. After processing, remove excess resin around the CM LOC® Housing of the female part with a round bur. Then finish and polish the prosthesis. Afterwards replace the processing insert in the CM LOC® Housing of the female part with a Pekkton® Retention insert in the desired force level.

 See description in Selection of retention inserts.



 Make sure that no polymer has flowed into the housing of the female part. If necessary, remove the processing insert and carefully remove excess polymer from the inside of the housing of the female part with a probe.



Selection of retention inserts

Four different CM LOC® Retention inserts made of Pekkton® are available for retention. The retention inserts are colour-coded and divided into four different levels of retention force.

yellow: extra low red: low
green: medium blue: strong

 Ensure that the selection of pull-off forces is adapted to the clinical situation. Only use the extra-low insert for immediate loading at the start.

 To provide the patient with comfortable and easy integration of the prosthesis as well as familiarisation with the retention in the mouth, it is recommended to fit the prosthesis with a CM LOC® Retention insert, extra-low, first. If the patient demands stronger retention, CM LOC® Retention inserts offering greater retention force can be used. For assembly and disassembly of the retention inserts, see description: assembly and disassembly of retention inserts.

Direct method: processing of the CM LOC® Housing during the treatment session.



It is essential to create sufficient space in the prosthesis prior to inclusion in the prosthetic body. Use a standard round bur for this purpose. There must not be any contact between the prosthesis and the CM LOC® Housing of the female part.



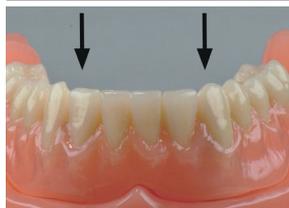
Mount the CM LOC® Block-out spacer on the male part.

 Make sure that the CM LOC® Block-out spacer fits well. The height of the CM LOC® Block-out spacer can be shortened to achieve better mounting of the female part.



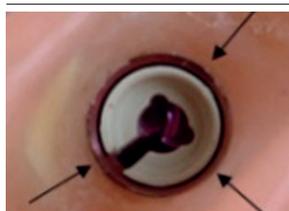
Then mount the CM LOC® Housing of the female part with mounted processing insert on the male part.

 Make sure that all undercuts are blocked out before polymerisation. Use a cold-curing polymer (e.g. GC Reline™, GC Advanced Technologies® Inc.) to anchor the CM LOC® Housing of the female part in the prosthesis. Apply the cold-curing polymer in the exposed area in the prosthesis and around the CM LOC® Housing of the female part.



Finalisation

Place the prosthesis on the CM LOC® Male part in the oral cavity. Make sure that the prosthesis is entirely in occlusion with the opposing jaw. Ensure that the prosthesis is retained passively without compression on the soft tissue while the cold-curing polymer cures. Excessive occlusal pressure during curing can cause the soft tissue to be compressed and then decompressed again. This can cause the processing inserts to then click out of position.



After processing, take the CM LOC® Block-out spacer out of the mouth.
 After processing, remove excess resin around the CM LOC® Housing of the female part with a round bur.
 Then finish and polish the prosthesis.
 Afterwards replace the processing insert in the CM LOC® Housing of the female part with a Pekkton® Retention insert in the desired force level.

-  See description in Selection of retention inserts.
-  Make sure that no polymer has flowed into the housing of the female part. If necessary, remove the processing insert and carefully remove excess polymer from the inside of the housing of the female part with a probe.



Selection of retention inserts

Four different CM LOC® Retention inserts made of Pekkton® are available for retention. The retention inserts are colour-coded and divided into four different levels of retention force.
 yellow: extra low red: low
 green: medium blue: strong

-  Ensure that the selection of pull-off forces is adapted to the clinical situation. Only use the extra-low insert for immediate loading at the start.
-  To provide the patient with comfortable and easy integration of the prosthesis as well as familiarisation with the retention in the mouth, it is recommended to fit the prosthesis with a CM LOC® Retention insert, extra-low, first. If the patient demands stronger retention, CM LOC® Retention inserts offering greater retention force can be used. For assembly and disassembly of the retention inserts, see description: assembly and disassembly of retention inserts.

Modifying an existing prosthesis using CM LOC® Components with simultaneous relining.



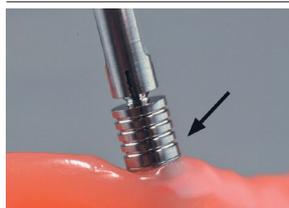
Remove existing anchorage from the patient's mouth.
 Use the CM LOC® Case Guide to determine the divergence of the implant axes between the implants. Place the CM LOC® Case Guide on the implant for this purpose by screwing in manually.

-  A corresponding CM LOC® Case Guide is available for each implant system.



By cyclically tipping, use the CM LOC® Case Guide to determine the implant axes, so that the individual implant axes to each other can be determined. Attention: view from side and front.

-  Should it not be possible to align the CM LOC® Case Guides parallel, a divergence of 20° between the implants is exceeded.
 If the divergence is greater than 20°, the CM LOC® Abutment may not be used, instead the CM LOC® FLEX is used.



Determining the abutment height

Choose the abutment height based on the implant position/gingival height and read off based on the graduation marks on the CM LOC® Case Guide. Determine the correct height of the CM LOC® Abutment with the lower edge of the CM LOC® Abutment positioned at least 1 mm above the gingiva. The lowest height starts at graduation marking 1.



Insert CM LOC® Abutment

First, push the CM LOC® Abutment from the side into the CM LOC® Screw Driver and screw it into the implant by hand.



Use the torque ratchet to tighten to the required torque. Make sure that the CM LOC® Screwdriver is correctly seated on the abutment. Secure all parts against aspiration.
 After assembly, the CM LOC® Screwdriver can be removed laterally by turning it back slightly.

-  The screwdriver features an ISO connection and fits onto the coupling inserts for the corresponding torque wrenches.



Relining

The previously mounted CM LOC® Housing of the female part with a mounted processing insert secures the prosthesis during impression-taking.



Mount the CM LOC® Block-out spacer on the abutment.

-  Make sure that the CM LOC® Block-out spacer fits well. The height of the CM LOC® Block-out spacer can be shortened when using different abutment heights to achieve better mounting of the female part. Make sure that all undercuts are blocked out before relining. After processing, take the CM LOC® Block-out spacer out of the mouth.

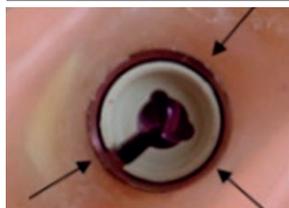


An impression of the relining with the existing prosthesis is then taken in the usual manner.

-  Do not apply impression material into the CM LOC® Housing for female parts and make sure that the prosthesis is securely seated on the CM LOC® Abutment. Otherwise, clean the CM LOC® Housing for female parts immediately.



The impression is then sent to the dental laboratory for fabrication of the model for relining using the conventional technique, as well as subsequent finishing and polishing of the prosthesis. Remove excess resin around the CM LOC® Housing of the female part with a round bur. Afterwards replace the processing insert in the CM LOC® Housing of the female part with a Pekkton® Retention insert in the desired force level.



-  See description in Selection of retention inserts.
-  Make sure that no polymer has flowed into the housing of the female part. If necessary, remove the processing insert and carefully remove excess polymer from the inside of the housing of the female part with a probe.



Selection of retention inserts

Four different CM LOC® Retention inserts made of Pekkton® are available for retention. The retention inserts are colour-coded and divided into four different levels of retention force.

yellow: extra low red: low
green: medium blue: strong

-  Ensure that the selection of pull-off forces is adapted to the clinical situation. Only use the extra-low insert for immediate loading at the start.
-  To provide the patient with comfortable and easy integration of the prosthesis as well as familiarisation with the retention in the mouth, it is recommended to fit the prosthesis with a CM LOC® Retention insert, extra-low, first. If the patient demands stronger retention, CM LOC® Retention inserts offering greater retention force can be used. For assembly and disassembly of the retention inserts, see description: assembly and disassembly of retention inserts.

The CM LOC® CAD/CAM Retention element as an additional retention element on a milled bar.



Conventional or digital impression of the clinical situation in the mouth and as specified by the implant manufacturer.

Subsequent fabrication of the master model in the laboratory.

The bar is then fabricated using CAD/CAM technology.

Please follow the manufacturer's instructions of the respective system.



-  When modelling the bar in the CAD software, allow for the position of the CM LOC® CAD/CAM Retention element. A standard thread M2 is required for fixation of the CM LOC® CAD/CAM Retention element in the CAD/CAM dental bar.



Insertion of the CM LOC® CAD/CAM Retention element

After the CAD/CAM dental bar has been fabricated, the CM LOC®CAD / CAM Retention element can now be mounted on the milled bar with the CM LOC® Screwdriver.

First, push the CM LOC® Abutment from the side into the CM LOC® Screwdriver and screw it into the bar by hand.

Use the torque ratchet to tighten to the required torque. Make sure that the CM LOC® Screwdriver is correctly seated on the abutment. Secure all parts against aspiration.

After assembly, the CM LOC® Screwdriver can be removed laterally by turning it back slightly.



 Torque of the CM LOC® CAD/CAM Retention element >35 Ncm.

 The screwdriver features an ISO connection and fits onto the coupling inserts for the corresponding torque wrenches.



After mounting of the milled bar with mounted CM LOC® CAD/CAM Retention element and fixed housing of the female part on the master cast, fabricate the denture in the conventional wax-up. Subsequent try-in of the wax-up on the patient.



Finalisation

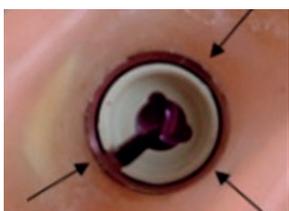
The prosthesis can now be fabricated using conventional techniques.

After processing, remove excess resin around the CM LOC® Housing of the female part with a round bur.

Then finish and polish the prosthesis.

Afterwards replace the processing insert in the CM LOC® Housing of the female part with a Pekkton® Retention insert in the desired force level.

 See description in Selection of retention inserts.



 Make sure that no polymer has flowed into the housing of the female part. If necessary, remove the processing insert and carefully remove excess polymer from the inside of the housing of the female part with a probe.



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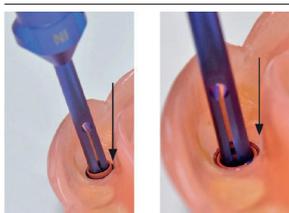
Assembly and disassembly of the retention inserts.



Assembly

The retention inserts are placed in the housing of the female part using the CM LOC® Multi-tool provided. Pick up the CM LOC® Retention insert with the IN side.





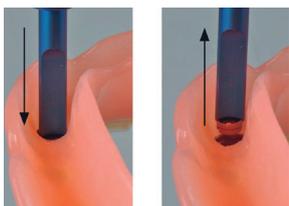
You can feel and hear the CM LOC® Retention insert lock into place. Press the CM LOC® Retention insert into the CM LOC® Housing of the female part in straight and parallel fashion until you can feel and hear it click into place.



Disassembly

Position the CM LOC® Multi-tool with the OUT side straight and parallel in the space between the CM LOC® Housing of the female part and the retention insert and press lightly into the CM LOC® Housing of the female part.

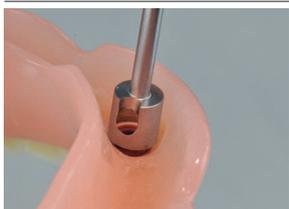
The CM LOC® Retention insert thus unlocks and can be removed in a straight manner from the CM LOC® Housing of the female part.



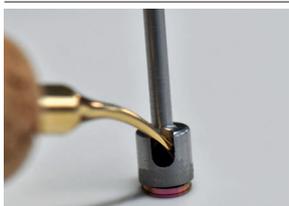
Disassembly CM LOC® Housing of the female part.



To disassemble the CM LOC® Housing of the female part, use the CM LOC® Extractor for the housing of the female part.



Mill the complete CM LOC® Housing of the female part with the CM LOC® Extractor for the housing of the female part.



Then remove the CM LOC® housing of the female part from the CM LOC® Extractor for the housing of the female part through the side opening with an instrument. For better withdrawal, it is recommended to briefly warm the CM LOC® Extractor for the housing of the female part over a flame.

19 Materials

More detailed information on the materials as well as their compositions can be found in the product-specific material data sheets, the product information as well as the product list compiled in Section 29. All relevant documents can be found on the website www.cmsa.ch/docs by entering the relevant product name.

20 Notes on storage



Insofar as no specific information on storage is given on the packaging of the product, we recommend storing the product in its original packaging, in a dry place, at room temperature and without direct sunlight. Improper storage can influence the product properties and lead to failure of the restoration.

21 Patient information

21.1 Handling / follow-up

On the day of insertion of the dentures at the latest, the patient must be informed that regular follow-up care is necessary to maintain the health of the entire masticatory system and the functionality of the denture. Ensure that the patients are motivated and instructed with regard to caring for their teeth as well as dentures.

Permanent and removable dentures are subject to considerable stress. Signs of wear are normal and cannot be avoided, only reduced. The amount of wear depends on the overall system.

Our endeavours are aimed at using materials that are as optimally matched as possible in order to reduce wear to an absolute minimum. Proper seating of the dentures on the mucosa must be checked at least once each year, and relining must be performed if required to prevent rocking movement (overload). We recommend checking the dentures at intervals of approx. 3 months initially and to replace the auxiliary parts such as retention inserts if necessary.

21.2 Insertion and removal of the dentures

It should be ensured that the dentures do not tilt, as any tilting can lead to damage. The denture should never be inserted by clenching the teeth, as this can damage or even break the connecting element.

Insertion

The denture can be placed on the anchor elements in the mouth using the thumb and index finger. Then it is correctly positioned on the anchoring elements applying gentle, even pressure. By carefully closing the jaws, it is possible to check whether the denture is in its correct final position.

Removal

For removal, the denture can be grasped with the thumb and index finger and carefully pulled from the anchor elements and taken out of the mouth.

21.3 Cleaning and care

We recommend cleaning teeth and dentures after every meal. Cleaning of dentures includes cleaning of the connecting element. Gentlest cleaning can be achieved by cleaning the restoration under running water with a soft toothbrush and the connecting element in the mouth with an interdental brush. The most intensive cleaning of the restoration is achieved with the aid of an ultrasonic device and a cleaning additive suitable for dentures.

Never clean the high precision connecting elements with toothpaste as this could lead to damage. Caution should also be exercised in the case of aggressive cleaning agents or tablets as this could damage the high-quality connecting element or impair its function.

Regular cleaning of the anchorage can prevent inflammation of the soft tissue.

22 Ordering information

The information relevant to your order can be found in the product list in Section 29 of this document. The product information is also helpful. This and other relevant documents can be found on the website www.cmsa.ch/docs by entering the relevant product name.

23 Availability

Some of the products described in this document may possibly not be available in all countries.

24 Traceability of the lot number

The lot numbers of all parts used must be documented to ensure traceability.

25 Complaint

Cendres+Métaux SA must be notified immediately of any incident that has occurred with regard to the product. To do this, please contact your customer advisor or send us your message by e-mail to the address complaints-cmbrand@cmsa.ch. In serious cases, also send a report to the competent authority where you are domiciled.

26 Safe disposal

The products must be disposed of in accordance with local laws and environmental regulations, taking into account the level of contamination. Cendres+Métaux Lux SA would be very pleased to accept precious metal waste. For information and additional details, please contact your Cendres+Métaux SA representative.

27 Trademarks

Registered trademarks of Cendres+Métaux Holding SA, Biel/Bienne, Switzerland include:

CM LOC®, Pekkton®

Unless explained specifically, all products marked with "®" are not registered trademarks of Cendres+Métaux Holding SA, but registered trademarks of the respective manufacturer.

28 Disclaimer

The manufacturer rejects any liability for damages resulting from non-compliance with these Instructions for Use. Cendres+Métaux SA products are parts of an overall concept and may only be used or combined with the appropriate original components and instruments. Otherwise, the manufacturer rejects any responsibility and liability. In case of complaints, please always include the lot number.

The use of third party products not distributed by Cendres+Métaux SA in connection with the products mentioned in the product list in Section 29 will void any warranty or other express or implied obligation of Cendres+Métaux SA.

Responsibility regarding the suitability of a product for the specific patient case is at the discretion of the specialist.

Cendres+Métaux SA disclaims any express or implied liability and shall not be responsible for any direct, indirect, punitive or other damages arising from or in connection with errors in professional judgement or practice in the use of Cendres+Métaux SA products.

The specialist is obliged to regularly study the latest developments of the products mentioned in the product list in Section 29 and their applications.

It should be noted that the descriptions contained in this document are not sufficient for the immediate application of Cendres+Métaux SA products. Expertise in dentistry, dental technology and instructions by an experienced specialist in the use of the products mentioned in the product list under Section 29 is always necessary.

In case of inconsistencies in translations, the English language version shall prevail.

29 Product list

Cat. No.	Product name	Material	Single use	Labelling	Basic UDI-DI
See Implant system	CM LOC® Abutment	TiAl6 V4 ELI, (Grade5)	Yes	CE 0483	764016651000045E5
05001304	CM LOC® CAD/CAM Retention Element for Bar	TiAl6 V4 ELI, (Grade5)	Yes	CE 0483	764016651000050DW
See Implant system	CM LOC® Case Guide	TiAl6 V4 ELI, (Grade5)	Yes	CE	764016651000056EA
05003001	CM LOC® Basic Set Titanium	TiAl6 V4 ELI, (Grade5) / Pekkton® / Santoprene	Yes	CE 0483	764016651000057EC
05001995	CM LOC® Housing Titanium for Pekkton® Inserts	TiAl6 V4 ELI, (Grade5)	Yes	CE 0483	764016651000053E4
05001314	CM LOC® Retention insert, extra-low	Pekkton®	Yes	CE 0483	764016651000053E4
05001315	CM LOC® Retention insert, low	Pekkton®	Yes	CE 0483	764016651000053E4
05001316	CM LOC® Retention insert, medium	Pekkton®	Yes	CE 0483	764016651000053E4
05001317	CM LOC® Retention insert, strong	Pekkton®	Yes	CE 0483	764016651000053E4
05001328	CM LOC® Processing insert	Pekkton®	Yes	CE	764016651000007DV
05001306	CM LOC® Housing Pekkton® for Pekkton® Inserts	Pekkton®	Yes	CE 0483	764016651000053E4
07000201	CM LOC® Spacer	Pekkton®	Yes	CE	764016651000026DZ
07000202	CM LOC® Block-out spacer	Santoprene	Yes	CE	764016651000027E3
07000204	CM LOC® Analog	TiAl6 V4 ELI, (Grade5)	Yes	CE	764016651000034DY
07000205	CM LOC® Multi-Tool for Pekkton® Retention insert	TiAl6 V4 ELI, (Grade5)	No	CE	764016651000001DH
07000206	CM LOC® Screw Driver	TiAl6 V4 ELI, (Grade5)	No	CE	764016651000022DR
07000213	CM LOC® Impression part	Pekkton®	Yes	CE	764016651000017DY
07000217	CM LOC® Housing Extractor	Steel	No	CE	764016651000009DZ
07000200	CM LOC® Instrument set	n/a	No	CE	764016651000025DX

Implant system

Cat. No.	Product name	Platform	Torque (recommendation)	Basic UDI-DI
Straumann				
0500 1408	CM LOC® Abutment	Straumann® RN 4.8 GH1	35Ncm	764016651000045E5
0500 1409	CM LOC® Abutment	Straumann® RN 4.8 GH2	35Ncm	764016651000045E5
0500 1410	CM LOC® Abutment	Straumann® RN 4.8 GH3	35Ncm	764016651000045E5
0500 1411	CM LOC® Abutment	Straumann® RN 4.8 GH4	35Ncm	764016651000045E5
0500 1412	CM LOC® Abutment	Straumann® RN 4.8 GH5	35Ncm	764016651000045E5
0500 1578	CM LOC® Case Guide	Straumann® RN 4.8	–	764016651000056EA
0500 1413	CM LOC® Abutment	Straumann® RC 4.1 / 4.8 GH1	35Ncm	764016651000045E5
0500 1414	CM LOC® Abutment	Straumann® RC 4.1 / 4.8 GH2	35Ncm	764016651000045E5
0500 1415	CM LOC® Abutment	Straumann® RC 4.1 / 4.8 GH3	35Ncm	764016651000045E5
0500 1416	CM LOC® Abutment	Straumann® RC 4.1 / 4.8 GH4	35Ncm	764016651000045E5
0500 1417	CM LOC® Abutment	Straumann® RC 4.1 / 4.8 GH5	35Ncm	764016651000045E5
0500 1579	CM LOC® Case Guide	Straumann® RC 4.1 / 4.8	–	764016651000056EA
0500 1418	CM LOC® Abutment	Straumann® NNC 3.5 GH1	35Ncm	764016651000045E5
0500 1419	CM LOC® Abutment	Straumann® NNC 3.5 GH2	35Ncm	764016651000045E5
0500 1420	CM LOC® Abutment	Straumann® NNC 3.5 GH3	35Ncm	764016651000045E5
0500 1421	CM LOC® Abutment	Straumann® NNC 3.5 GH4	35Ncm	764016651000045E5
0500 1422	CM LOC® Abutment	Straumann® NNC 3.5 GH5	35Ncm	764016651000045E5
0500 1580	CM LOC® Case Guide	Straumann® NNC 3.5	–	764016651000056EA
0500 2547	CM LOC® Abutment	Straumann® NC 3.3 GH1	35Ncm	764016651000045E5
0500 2548	CM LOC® Abutment	Straumann® NC 3.3 GH2	35Ncm	764016651000045E5
0500 2549	CM LOC® Abutment	Straumann® NC 3.3 GH3	35Ncm	764016651000045E5
0500 2550	CM LOC® Abutment	Straumann® NC 3.3 GH4	35Ncm	764016651000045E5
0500 2551	CM LOC® Abutment	Straumann® NC 3.3 GH5	35Ncm	764016651000045E5
0500 2552	CM LOC® Case Guide	Straumann® NC 3.3	–	764016651000056EA
Nobel Biocare				
0500 1423	CM LOC® Abutment	Nobel Biocare Replace Select® NP 3.5 GH1	35Ncm	764016651000045E5
0500 1424	CM LOC® Abutment	Nobel Biocare Replace Select® NP 3.5 GH2	35Ncm	764016651000045E5
0500 1425	CM LOC® Abutment	Nobel Biocare Replace Select® NP 3.5 GH3	35Ncm	764016651000045E5
0500 1426	CM LOC® Abutment	Nobel Biocare Replace Select® NP 3.5 GH4	35Ncm	764016651000045E5
0500 1427	CM LOC® Abutment	Nobel Biocare Replace Select® NP 3.5 GH5	35Ncm	764016651000045E5
0500 1581	CM LOC® Case Guide	Nobel Biocare Replace Select® NP 3.5	–	764016651000056EA
0500 1296	CM LOC® Abutment	Nobel Biocare Replace Select® RP 4.3 GH1	35Ncm	764016651000045E5
0500 1300	CM LOC® Abutment	Nobel Biocare Replace Select® RP 4.3 GH2	35Ncm	764016651000045E5

Cat. No.	Product name	Platform	Torque (recom- mendation)	Basic UDI-DI
0500 1301	CM LOC® Abutment	Nobel Biocare Replace Select® RP 4.3 GH3	35Ncm	764016651000045E5
0500 1302	CM LOC® Abutment	Nobel Biocare Replace Select® RP 4.3 GH4	35Ncm	764016651000045E5
0500 1303	CM LOC® Abutment	Nobel Biocare Replace Select® RP 4.3 GH5	35Ncm	764016651000045E5
0500 1582	CM LOC® Case Guide	Nobel Biocare Replace Select® RP 4.3	–	764016651000056EA
0500 2109	CM LOC® Abutment	Nobel Biocare Active® NP 3.5 GH1	35Ncm	764016651000045E5
0500 2110	CM LOC® Abutment	Nobel Biocare Active® NP 3.5 GH2	35Ncm	764016651000045E5
0500 2111	CM LOC® Abutment	Nobel Biocare Active® NP 3.5 GH3	35Ncm	764016651000045E5
0500 2112	CM LOC® Abutment	Nobel Biocare Active® NP 3.5 GH4	35Ncm	764016651000045E5
0500 2113	CM LOC® Abutment	Nobel Biocare Active® NP 3.5 GH5	35Ncm	764016651000045E5
0500 2187	CM LOC® Case Guide	Nobel Biocare Active® NP 3.5	–	764016651000056EA
0500 1437	CM LOC® Abutment	Nobel Biocare Active® RP 4.3/5.0 GH1	35Ncm	764016651000045E5
0500 1438	CM LOC® Abutment	Nobel Biocare Active® RP 4.3/5.0 GH2	35Ncm	764016651000045E5
0500 1439	CM LOC® Abutment	Nobel Biocare Active® RP 4.3/5.0 GH3	35Ncm	764016651000045E5
0500 1440	CM LOC® Abutment	Nobel Biocare Active® RP 4.3/5.0 GH4	35Ncm	764016651000045E5
0500 1441	CM LOC® Abutment	Nobel Biocare Active® RP 4.3/5.0 GH5	35Ncm	764016651000045E5
0500 1584	CM LOC® Case Guide	Nobel Biocare Active® RP 4.3/5.0	–	764016651000056EA
0500 2114	CM LOC® Abutment	Nobel Biocare Brånemark® RP 4.0 GH1	35Ncm	764016651000045E5
0500 2115	CM LOC® Abutment	Nobel Biocare Brånemark® RP 4.0 GH2	35Ncm	764016651000045E5
0500 2116	CM LOC® Abutment	Nobel Biocare Brånemark® RP 4.0 GH3	35Ncm	764016651000045E5
0500 2117	CM LOC® Abutment	Nobel Biocare Brånemark® RP 4.0 GH4	35Ncm	764016651000045E5
0500 2118	CM LOC® Abutment	Nobel Biocare Brånemark® RP 4.0 GH5	35Ncm	764016651000045E5
0500 2188	CM LOC® Case Guide	Nobel Biocare Brånemark® RP 4.0	–	764016651000056EA
Astra Tech				
0500 1452	CM LOC® Abutment	Astra Tech OsseoSpeed® 3.5/4.0 GH1	25Ncm	764016651000045E5
0500 1453	CM LOC® Abutment	Astra Tech OsseoSpeed® 3.5/4.0 GH2	25Ncm	764016651000045E5
0500 1454	CM LOC® Abutment	Astra Tech OsseoSpeed® 3.5/4.0 GH3	25Ncm	764016651000045E5
0500 1455	CM LOC® Abutment	Astra Tech OsseoSpeed® 3.5/4.0 GH4	25Ncm	764016651000045E5
0500 1456	CM LOC® Abutment	Astra Tech OsseoSpeed® 3.5/4.0 GH5	25Ncm	764016651000045E5
0500 1587	CM LOC® Case Guide	Astra Tech OsseoSpeed® 3.5/4.0	–	764016651000056EA
0500 1457	CM LOC® Abutment	Astra Tech OsseoSpeed® 4.5/5.0 GH1	25Ncm	764016651000045E5
0500 1458	CM LOC® Abutment	Astra Tech OsseoSpeed® 4.5/5.0 GH2	25Ncm	764016651000045E5
0500 1459	CM LOC® Abutment	Astra Tech OsseoSpeed® 4.5/5.0 GH3	25Ncm	764016651000045E5
0500 1460	CM LOC® Abutment	Astra Tech OsseoSpeed® 4.5/5.0 GH4	25Ncm	764016651000045E5
0500 1461	CM LOC® Abutment	Astra Tech OsseoSpeed® 4.5/5.0 GH5	25Ncm	764016651000045E5
0500 1588	CM LOC® Case Guide	Astra Tech OsseoSpeed® 4.5/5.0	–	764016651000056EA
0500 1963	CM LOC® Abutment	Astra Tech OsseoSpeed® EV 3.6 GH1	25Ncm	764016651000045E5
0500 1964	CM LOC® Abutment	Astra Tech OsseoSpeed® EV 3.6 GH2	25Ncm	764016651000045E5
0500 1965	CM LOC® Abutment	Astra Tech OsseoSpeed® EV 3.6 GH3	25Ncm	764016651000045E5
0500 1966	CM LOC® Abutment	Astra Tech OsseoSpeed® EV 3.6 GH4	25Ncm	764016651000045E5
0500 1967	CM LOC® Abutment	Astra Tech OsseoSpeed® EV 3.6 GH5	25Ncm	764016651000045E5
0500 1981	CM LOC® Case Guide	Astra Tech OsseoSpeed® EV 3.6	–	764016651000056EA
0500 1968	CM LOC® Abutment	Astra Tech OsseoSpeed® EV 4.2 GH1	25Ncm	764016651000045E5
0500 1969	CM LOC® Abutment	Astra Tech OsseoSpeed® EV 4.2 GH2	25Ncm	764016651000045E5
0500 1970	CM LOC® Abutment	Astra Tech OsseoSpeed® EV 4.2 GH3	25Ncm	764016651000045E5
0500 1971	CM LOC® Abutment	Astra Tech OsseoSpeed® EV 4.2 GH4	25Ncm	764016651000045E5
0500 1972	CM LOC® Abutment	Astra Tech OsseoSpeed® EV 4.2 GH5	25Ncm	764016651000045E5
0500 1982	CM LOC® Case Guide	Astra Tech OsseoSpeed® EV 4.2	–	764016651000056EA
0500 1973	CM LOC® Abutment	Astra Tech OsseoSpeed® EV 4.8 GH1	25Ncm	764016651000045E5
0500 1974	CM LOC® Abutment	Astra Tech OsseoSpeed® EV 4.8 GH2	25Ncm	764016651000045E5
0500 1975	CM LOC® Abutment	Astra Tech OsseoSpeed® EV 4.8 GH3	25Ncm	764016651000045E5
0500 1976	CM LOC® Abutment	Astra Tech OsseoSpeed® EV 4.8 GH4	25Ncm	764016651000045E5
0500 1977	CM LOC® Abutment	Astra Tech OsseoSpeed® EV 4.8 GH5	25Ncm	764016651000045E5
0500 1983	CM LOC® Case Guide	Astra Tech OsseoSpeed® EV 4.8	–	764016651000056EA
Zimmer				
0500 1462	CM LOC® Abutment	Zimmer Tapered Screw-Vent® 3.5 GH1	30Ncm	764016651000045E5
0500 1463	CM LOC® Abutment	Zimmer Tapered Screw-Vent® 3.5 GH2	30Ncm	764016651000045E5
0500 1464	CM LOC® Abutment	Zimmer Tapered Screw-Vent® 3.5 GH3	30Ncm	764016651000045E5
0500 1465	CM LOC® Abutment	Zimmer Tapered Screw-Vent® 3.5 GH4	30Ncm	764016651000045E5
0500 1466	CM LOC® Abutment	Zimmer Tapered Screw-Vent® 3.5 GH5	30Ncm	764016651000045E5
0500 1589	CM LOC® Case Guide	Zimmer Tapered Screw-Vent® 3.5	–	764016651000056EA
0500 1467	CM LOC® Abutment	Zimmer Tapered Screw-Vent® 4.5 GH1	30Ncm	764016651000045E5
0500 1468	CM LOC® Abutment	Zimmer Tapered Screw-Vent® 4.5 GH2	30Ncm	764016651000045E5
0500 1469	CM LOC® Abutment	Zimmer Tapered Screw-Vent® 4.5 GH3	30Ncm	764016651000045E5
0500 1470	CM LOC® Abutment	Zimmer Tapered Screw-Vent® 4.5 GH4	30Ncm	764016651000045E5
0500 1471	CM LOC® Abutment	Zimmer Tapered Screw-Vent® 4.5 GH5	30Ncm	764016651000045E5
0500 1590	CM LOC® Case Guide	Zimmer Tapered Screw-Vent® 4.5	–	764016651000056EA

Cat. No.	Product name	Platform	Torque (recom- mendation)	Basic UDI-DI
MIS (standard narrow)				
0500 1462	CM LOC® Abutment	MiS® Seven 3.5 GH1	30Ncm	764016651000045E5
0500 1463	CM LOC® Abutment	MiS® Seven 3.5 GH2	30Ncm	764016651000045E5
0500 1464	CM LOC® Abutment	MiS® Seven 3.5 GH3	30Ncm	764016651000045E5
0500 1465	CM LOC® Abutment	MiS® Seven 3.5 GH4	30Ncm	764016651000045E5
0500 1466	CM LOC® Abutment	MiS® Seven 3.5 GH5	30Ncm	764016651000045E5
0500 1589	CM LOC® Case Guide	MiS® Seven 3.5	–	764016651000056EA
MIS (wide platform)				
0500 1467	CM LOC® Abutment	MiS® Seven 4.5 GH1	30Ncm	764016651000045E5
0500 1468	CM LOC® Abutment	MiS® Seven 4.5 GH2	30Ncm	764016651000045E5
0500 1469	CM LOC® Abutment	MiS® Seven 4.5 GH3	30Ncm	764016651000045E5
0500 1470	CM LOC® Abutment	MiS® Seven 4.5 GH4	30Ncm	764016651000045E5
0500 1471	CM LOC® Abutment	MiS® Seven 4.5 GH5	30Ncm	764016651000045E5
0500 1590	CM LOC® Case Guide	MiS® Seven 4.5	–	764016651000056EA
BioHorizons				
0500 1462	CM LOC® Abutment	BioHorizons® Internal 3.5 GH1	30Ncm	764016651000045E5
0500 1463	CM LOC® Abutment	BioHorizons® Internal 3.5 GH2	30Ncm	764016651000045E5
0500 1464	CM LOC® Abutment	BioHorizons® Internal 3.5 GH3	30Ncm	764016651000045E5
0500 1465	CM LOC® Abutment	BioHorizons® Internal 3.5 GH4	30Ncm	764016651000045E5
0500 1466	CM LOC® Abutment	BioHorizons® Internal 3.5 GH5	30Ncm	764016651000045E5
0500 1589	CM LOC® Case Guide	BioHorizons® Internal 3.5	–	764016651000056EA
0500 1467	CM LOC® Abutment	BioHorizons® Internal 4.5 GH1	30Ncm	764016651000045E5
0500 1468	CM LOC® Abutment	BioHorizons® Internal 4.5 GH2	30Ncm	764016651000045E5
0500 1469	CM LOC® Abutment	BioHorizons® Internal 4.5 GH3	30Ncm	764016651000045E5
0500 1470	CM LOC® Abutment	BioHorizons® Internal 4.5 GH4	30Ncm	764016651000045E5
0500 1471	CM LOC® Abutment	BioHorizons® Internal 4.5 GH5	30Ncm	764016651000045E5
0500 1590	CM LOC® Case Guide	BioHorizons® Internal 4.5	–	764016651000056EA
Camlog				
0500 1544	CM LOC® Abutment	Camlog® 3.8 GH1	30Ncm	764016651000045E5
0500 1545	CM LOC® Abutment	Camlog® 3.8 GH2	30Ncm	764016651000045E5
0500 1546	CM LOC® Abutment	Camlog® 3.8 GH3	30Ncm	764016651000045E5
0500 1547	CM LOC® Abutment	Camlog® 3.8 GH4	30Ncm	764016651000045E5
0500 1591	CM LOC® Case Guide	Camlog® 3.8	–	764016651000056EA
0500 1549	CM LOC® Abutment	Camlog® 4.3 GH1	30Ncm	764016651000045E5
0500 1550	CM LOC® Abutment	Camlog® 4.3 GH2	30Ncm	764016651000045E5
0500 1551	CM LOC® Abutment	Camlog® 4.3 GH3	30Ncm	764016651000045E5
0500 1552	CM LOC® Abutment	Camlog® 4.3 GH4	30Ncm	764016651000045E5
0500 1592	CM LOC® Case Guide	Camlog® 4.3	–	764016651000056EA
0500 1482	CM LOC® Abutment	Conelog® 3.8/4.3 GH1	30Ncm	764016651000045E5
0500 1483	CM LOC® Abutment	Conelog® 3.8/4.3 GH2	30Ncm	764016651000045E5
0500 1484	CM LOC® Abutment	Conelog® 3.8/4.3 GH3	30Ncm	764016651000045E5
0500 1485	CM LOC® Abutment	Conelog® 3.8/4.3 GH4	30Ncm	764016651000045E5
0500 1486	CM LOC® Abutment	Conelog® 3.8/4.3 GH5	30Ncm	764016651000045E5
0500 1594	CM LOC® Case Guide	Conelog® 3.8/4.3	–	764016651000056EA
Dentsply				
05002014	CM LOC® Abutment	Dentsply Ankylos® C 3.5, 4.5, 5.5, 7.0 GH1	25Ncm	764016651000045E5
05002015	CM LOC® Abutment	Dentsply Ankylos® C 3.5, 4.5, 5.5, 7.0 GH2	25Ncm	764016651000045E5
05002016	CM LOC® Abutment	Dentsply Ankylos® C 3.5, 4.5, 5.5, 7.0 GH3	25Ncm	764016651000045E5
05002017	CM LOC® Abutment	Dentsply Ankylos® C 3.5, 4.5, 5.5, 7.0 GH4	25Ncm	764016651000045E5
05002018	CM LOC® Abutment	Dentsply Ankylos® C 3.5, 4.5, 5.5, 7.0 GH5	25Ncm	764016651000045E5
05002020	CM LOC® Case Guide	Dentsply Ankylos® C 3.5, 4.5, 5.5, 7.0	–	764016651000056EA
Sweden+Martina				
05002119	CM LOC® Abutment	Sweden+Martina Premium Kohno 3.3 GH1	30Ncm	764016651000045E5
05002120	CM LOC® Abutment	Sweden+Martina Premium Kohno 3.3 GH2	30Ncm	764016651000045E5
05002121	CM LOC® Abutment	Sweden+Martina Premium Kohno 3.3 GH3	30Ncm	764016651000045E5
05002122	CM LOC® Abutment	Sweden+Martina Premium Kohno 3.3 GH4	30Ncm	764016651000045E5
05002123	CM LOC® Abutment	Sweden+Martina Premium Kohno 3.3 GH5	30Ncm	764016651000045E5
05002189	CM LOC® Case Guide	Sweden+Martina Premium Kohno 3.3	–	764016651000056EA
05002124	CM LOC® Abutment	Sweden+Martina Premium Kohno 3.8 GH1	30Ncm	764016651000045E5
05002125	CM LOC® Abutment	Sweden+Martina Premium Kohno 3.8 GH2	30Ncm	764016651000045E5
05002126	CM LOC® Abutment	Sweden+Martina Premium Kohno 3.8 GH3	30Ncm	764016651000045E5
05002127	CM LOC® Abutment	Sweden+Martina Premium Kohno 3.8 GH4	30Ncm	764016651000045E5
05002128	CM LOC® Abutment	Sweden+Martina Premium Kohno 3.8 GH5	30Ncm	764016651000045E5
05002190	CM LOC® Case Guide	Sweden+Martina Premium Kohno 3.8	–	764016651000056EA
05002129	CM LOC® Abutment	Sweden+Martina Premium Kohno 4.25 GH1	30Ncm	764016651000045E5
05002130	CM LOC® Abutment	Sweden+Martina Premium Kohno 4.25 GH2	30Ncm	764016651000045E5
05002131	CM LOC® Abutment	Sweden+Martina Premium Kohno 4.25 GH3	30Ncm	764016651000045E5

Cat. No.	Product name	Platform	Torque (recom- mendation)	Basic UDI-DI
05002132	CM LOC® Abutment	Sweden+Martina Premium Kohno 4.25 GH4	30Ncm	764016651000045E5
05002133	CM LOC® Abutment	Sweden+Martina Premium Kohno 4.25 GH5	30Ncm	764016651000045E5
05002192	CM LOC® Case Guide	Sweden+Martina Premium Kohno 4.25/5.0/6.0	–	764016651000056EA
05002134	CM LOC® Abutment	Sweden+Martina Premium Kohno 5.0/6.0 GH1	30Ncm	764016651000045E5
05002135	CM LOC® Abutment	Sweden+Martina Premium Kohno 5.0/6.0 GH2	30Ncm	764016651000045E5
05002136	CM LOC® Abutment	Sweden+Martina Premium Kohno 5.0/6.0 GH3	30Ncm	764016651000045E5
05002137	CM LOC® Abutment	Sweden+Martina Premium Kohno 5.0/6.0 GH4	30Ncm	764016651000045E5
05002138	CM LOC® Abutment	Sweden+Martina Premium Kohno 5.0/6.0 GH5	30Ncm	764016651000045E5
05002192	CM LOC® Case Guide	Sweden+Martina Premium Kohno 4.25/5.0/6.0	–	764016651000056EA
Osstem				
05002159	CM LOC® Abutment	Osstem® T _s Regular 4.0/4.5/5.0/6.0/7.0 GH1	30Ncm	764016651000045E5
05002160	CM LOC® Abutment	Osstem® T _s Regular 4.0/4.5/5.0/6.0/7.0 GH2	30Ncm	764016651000045E5
05002161	CM LOC® Abutment	Osstem® T _s Regular 4.0/4.5/5.0/6.0/7.0 GH3	30Ncm	764016651000045E5
05002162	CM LOC® Abutment	Osstem® T _s Regular 4.0/4.5/5.0/6.0/7.0 GH4	30Ncm	764016651000045E5
05002163	CM LOC® Abutment	Osstem® T _s Regular 4.0/4.5/5.0/6.0/7.0 GH5	30Ncm	764016651000045E5
05002197	CM LOC® Case Guide	Osstem® T _s Regular 4.0/4.5/5.0/6.0/7.0	–	764016651000056EA
05002590	CM LOC® Abutment	Osstem® T _s Mini 3.5 GH1	30Ncm	764016651000045E5
05002591	CM LOC® Abutment	Osstem® T _s Mini 3.5 GH2	30Ncm	764016651000045E5
05002592	CM LOC® Abutment	Osstem® T _s Mini 3.5 GH3	30Ncm	764016651000045E5
05002593	CM LOC® Abutment	Osstem® T _s Mini 3.5 GH4	30Ncm	764016651000045E5
05002594	CM LOC® Abutment	Osstem® T _s Mini 3.5 GH5	30Ncm	764016651000045E5
05002596	CM LOC® Case Guide	Osstem® T _s Mini 3.5	–	764016651000056EA
05002154	CM LOC® Abutment	Osstem® US Regular 4.1 GH1	30Ncm	764016651000045E5
05002155	CM LOC® Abutment	Osstem® US Regular 4.1 GH2	30Ncm	764016651000045E5
05002156	CM LOC® Abutment	Osstem® US Regular 4.1 GH3	30Ncm	764016651000045E5
05002157	CM LOC® Abutment	Osstem® US Regular 4.1 GH4	30Ncm	764016651000045E5
05002158	CM LOC® Abutment	Osstem® US Regular 4.1 GH5	30Ncm	764016651000045E5
05002196	CM LOC® Case Guide	Osstem® US Regular 4.1	–	764016651000056EA
05002149	CM LOC® Abutment	Osstem® SS Regular 4.8 GH1	30Ncm	764016651000045E5
05002150	CM LOC® Abutment	Osstem® SS Regular 4.8 GH2	30Ncm	764016651000045E5
05002151	CM LOC® Abutment	Osstem® SS Regular 4.8 GH3	30Ncm	764016651000045E5
05002152	CM LOC® Abutment	Osstem® SS Regular 4.8 GH4	30Ncm	764016651000045E5
05002153	CM LOC® Abutment	Osstem® SS Regular 4.8 GH5	30Ncm	764016651000045E5
05002195	CM LOC® Case Guide	Osstem® SS Regular 4.8	–	764016651000056EA

30 Labelling on packaging/symbols

	Date of manufacture
	Manufacturer
	Catalogue number
	Lot number
	Quantity
 www.cmsa.ch/docs	Observe the Instructions for Use, which are available in electronic form at the address specified.
Rx only	Attention: According to US federal law, this product may only be sold by or on behalf of a physician.
	Cendres+Métaux products with CE labelling meet the requirements of the relevant European requirements.
	Do not re-use
	Non-sterile
	Protect from sunlight
	Attention, observe accompanying documents
	Clear product identification
	European Authorised Representative
	Importer
	Medical device