Sleep disorders ... dentistry?

Ser the

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Sleep disorders ... dentistry?

Snoring and the obstructive sleep apnea (OSA), the interruption of the respiration during the sleep, belong to the most common sleep disorders.

In western industrialized countries approx. 10 % of the 20 years old men and 50 % of the 50 years old men snore. Women come up to two third of these data.

These sleep disorders can be successfully treated or at least be positively influenced in dentistry.



open respiratory tracts: Silensor-sl, mandibular advancement splint



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Topics of this lecture

- 1. Sleep disorders
- 2. Apnea during sleep
- 3. Therapy:
- 1. NCPAP (Nasal Continuous Positive Airways Pressure)
- 1. Surgery
- 2. Oral appliances
- 1. Tongue retainer
- 2. Protrusion splints (MAS, mandibular advancement splint) splints that advance the lower jaw

Terms on somnology:

Silensor-sl

primary snoring →	light snoring
rhonchopathy ->	upper airway resistance syndrom (UARS) = obstructive snoring
Apnea 🔶	pauses in breathing
obstructive apnea →	relocation of the upper respiratory tract
central apnea →	stoppage of the respiration reflex
Hypopnea 🔶	air flow reduced by more than 50 %
AHI 🔶	apnea/hypopnea index
RDI 🔿	respiratory disturbance index = disorder of the respiration
	reaction of awakening (no waking state)
somnography 🔿	examination of the sleep quality
SBAS ->	sleep-related breathing disorder





Sleep disorders

External sleep disorders

Disturbance of the sleep quality by external reasons, noise, bad bed, bad room air etc.

Internal sleep disorders

Disturbance of the sleep quality by internal reasons, teeth grinding (bruxism), indigestive food, medicaments, drugs, snoring, sleep apnea, pain and other diseases.





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Snoring

open pharynx, no noise

Snoring is generated in the upper respiratory tract.

By the decreasing of the muscle tone during sleep the lower jaw falls backwards.

The result is a narrowing of the respiratory tract.

The air flow will be accelerated and soft tissue (velum, uvula and other) start to vibrate and cause the snoring noise. Snoring is a mechanical process that can be counteracted mechanically. This is where the dental therapy comes into action.

Contrary to the rhonchopathy, the abnormal snoring, the light or primary snoring does not have a negative influence on the cardiovascular system and the oxygen supply.



narrowed pharynx, accelerated air flow at the same air volume, noise caused by flittering, vibrating structures





Snoring

	abnormal snoring	: harmless, primary snoring:
→ Abnormal snoring:		
	frequenc	y — frequency
Rhonchopathy, obstructive	snoring every nigh	t snoring from time to time
snoring, snoring with		
airway resistance.	loudness of snorin	
Upper airway resistance syndrom (UARS)	very loud, audible in the next roor	n moderately loud to loud, harmonious
	soun	d ——— sound
	explosive, hard, wit	
→ Primary snoring:	high frequency	• •
er er en er	stertorou	
Snoring harmless to health		
J	respiratio	n ——— respiration
	irregular, possibly with pause	•
	(breaks)
	sleep pattern	
	restless sleep, frequer	
	awakenings	



Commercial offers against primary snoring

On the market there are numerous devices against snoring available.

In case of very light snoring a success might be achieved but some products are rather questionable.

These auxiliaries are available to anyone but cannot always be considered as being harmless.



Auxiliaries to avoid a dorsial position.



schnarchladen.de

Nostril spreader, the reason for snoring, however, is mostly in the pharyngeal area.



Pillow Silensor (Canada), let the head tilt to the side.

(This pillow is the reason why the Erkodent Silensor is called Silent Nite in North America.)



spray to "grease" the mucosa

spray to stiffen the velum by foaming





nose inserts for stimulation









The apnea and hypopnea

The apnea is a complete respiratory stop. The obstructive apnea (OSA) is a mechanical relocation of the respiratory tract. In case of a central apnea the central respiration reflex stops.

The hypopnea is a reduction of the air flow by more than 50 % (reduced oxygen saturation.)

Index, AHI (RDI) is the degree for the severity of the disease.

An apnea/hypopnea lasts at least 10 sec. The apnea per hour sleep are counted, 10 apnea per hour result in an index of 10.

An index of 0 to 5 is normal, when the index is 5 to 10 there is a median disease and at an index of 20 and more there is a severe disease. In other countries the graduation is slightly different.

An obstructive apnea is characterized by an interruption of the very noiseful rhonchopathy. A central apnea runs without external signs.



obstructive sleep apnea, obstructed pharynx area of the obstruction





Symptoms and consequences of the OSA

The main symptom of the obstructive sleep apnea is a loud, irregular snoring, the rhonchopathy.

Family members might also report of pauses in breathing that are ended by the "arousal", a loud "implosive" restart of the snoring.

A non-treated OSA leads to mostly chronical cardiovascular diseases.

Further symptoms of the OSA:

- → restless sleep with difficulties staying asleep
- → tiredness during the day, microsleep, strong desire to sleep
- → headache and vertigo after awakening and getting up
- \rightarrow dry mouth
- → sweating during the night
- → increased urge to urinate during the night (nycturia)
- → lack of concentration
- → depressive mood
- → impotence (erectile dysfunction)

Health consequences of the OSA:

- → hypertension
- → myocardial infarction
- → apoplectic stroke
- → sudden cardiac death with increased probability
- → cardiac arrhythmia
- → depressions
- → stomach ulcer, hearing loss and other stress diseases
- → diabetes mellitus



Diagnostic investigation:

Ambulatory snoring investigation: Home screening, mostly an investigation for one night at home.

In-patient polysomnography: In a sleep laboratory, mostly for two nights in order to minimize influences of the unfamiliar surroundings. Diagnostic apparatus usable at home, in familiar surroundings, for collection of:

- → air-flow and snoring
- → heart rate
- → oxygen saturation
- → body position
- → thoracic and abdominal movements

Extensive investigation in a sleep laboratory for collection of:

- → electro-encephalogram of the brain (EEG)
- → rhythm of the heart (ECG)
- \rightarrow oxygen level of the blood (pulse oxymetry)
- → body temperature
- → air-flow (mouth and nose)
- → respiratory movement
- → muscle tension (EMG)
- → leg movement
- → eye movement (EOG)
- → body position









Therapy

(N) CPAP (Nasal) Continuous Positive Air Pressure.

Continuous (nasal) positive pressure of the respiratory air during the sleep.

The NCPAP respiratory mask is the therapeutical "gold standard" in case of an obstructive sleep apnea.

The side effects like irritated nasal mucosa and pharynx by the desiccative air flow, irritated conjunctiva caused by leakages, pressure points and skin irritations, however, lead to a low acceptance (compliance), especially in case of a not severe disease.



NCPAP function: Opening of the pharynx by continuous positive pressure of the respiratory air.

Modern units adapt the pressure to the respective airway resistance, also the Cheyne-Stokes-respiration is included.

The Cheyne-Stokes-respiration often occurs during the night at patients with advanced cardiac insufficiency (central sleep apnea syndrome). The result is a periodically recurring rise and fall of the respiration with changing respiratory frequence and respiratory pauses.









Therapy

Surgery: surgical correction of a retrognathism. Complex, effective surgery. In very rare cases executed because of an OSA.

UPPP (Uvula-Palato-Pharyngo-Plastic)

Laser- and radio frequency ablation:

Cicacitration to tighten the soft palate and correction of the uvula and the ending soft palate.

(for radio frequency ablation please see http://www.dr aschmann.de/de/behandlung.htm)

Implantive surgery, plastic rods are implanted to stiffen the structures.



Illustrations Dr. Pelosi, Parma

The correction of the uvula and the ending soft palate is acceptable. The effectiveness, however, is evaluated very differently.







The total removal of the uvula has to be seen very critically because of the very inconvenient side-effects.







Oral appliances

tongue retainer keep the tongue in an advanced position

protrusion splints resp. MAS (mandibular advancement splints) keep the lower jaw in a defined advancement

Protrusion splints that are realized as monobloc allow no or only little movement of the lower jaw. Studies have found that for this reason, there are slightly more TMJ (temporo mandibular joint) disorders to expect.

Therefore monobloc constructions should be rejected.







Aveo TSD, also suitable for edentulous persons

Lower jaw and tongue are connected to each other in the area of the frontal pharynx by ligaments and muscles. An advancement of these structures increases the passage of the pharynx and thus reduces the obstruction.

MAS can function by pulling:



TAP-T

or by pushing:







Sommnodent

IST, modified Herbst device: Attention, If the mouth falls open, the respiratory tract is additionally narrowed. The device is thereby counter- productive, that means the mouth opening has to be limited (arrow)!





Protrusion splints

The effectiveness of protrusion splints or MAS (mandibular advancement splints) is based on the fxation or defined advancement of the lower jaw.

Protrusion splints have a higher acceptance than tongue retainers, provided that they are not very voluminous and allow jaw movements.

If the lower jaw is advanced by 4 mm the front wall of the pharynx moves by about half to the front.

Snoring, narrowed pharynx

> By advancement of the lower jaw and the root of the tongue the protrusion splint opens the respiratory tract in the pharyngeal area.

obstructive sleep apnea, closed pharynx





Protrusion splints

Constructions based on pulling:

Silensor-sl

Specially shaped connectors keep the lower jaw in a certain advancement.

The connectors are easily replaceable. The Silensor-sl is adjustable in graduations of 1 mm.

The Silensor-sl allows relatively large jaw movements.

The construction is metal-free to avoid galvanic currents.



The versions that are based on pulling allow a large mouth opening and thereby additionally enlarge the respiratory tract.

Silensor-sl:

In case of sudden propulsion movements the anchors may glide in the connectors.

The S-shape avoids a hard stop of the connectors.











Thermoforming

Protrusion splints

Silensor-sl:

Therapeutical efficiency

X-ray examination of the University in Parma.

Advancement Silensor-sl: 4 mm









Protrusion splints

Therapy with protrusion splints:

To avoid failures and to minimize undesired side effects there are some conditions and requirements to be observed.

The dentist should have experience in the splint therapy, evaluate the oral situation and know possible side effects.

On suspicion of an apnea disease the patient should be transferred to specialists.

The patient:

Contraindications for the therapy with the Silensor-sl:

Apnea-index higher than 20 (more than 20 pauses in respiration per hour during sleep).

Inflammatory, painful temporomandibular jaw problems.

Loose tooth anchorage.

Less than 8 teeth per jaw.

Prognathic bite, protruding lower jaw, an advancement of the lower jaw is mostly not possible.

BMI (Body-Mass-Index) higher than 30. If the BMI is higher, the effect decreases continuously.

(BMI= body weight (kg) divided by body size2 (m))

The same applies if the neck size exceeds 44 cm.

The splint

should fulfill the following requirements:

All teeth are included.

No stiff connection of the jaws.

Individual adaptation with firm fit.

The advancement is adjustable.

Because of the needed acceptance the appliance is as comfortable as possible.

Metal free constructions are to be preferred.



Protrusion splints

Therapy with protrusion splints,

the side effects:

The most important side effect are <u>tooth movements</u>, the splint should therefore compass all teeth, a firm attachment apparatus is required. Self control and control by the dentist is necessary.

Unpleasant, dragging sensation in the joint area that, however, decreases with the time.

<u>Morning malocclusion</u>, after removal of the splint the patient feels a more or less considerably changed bite situation until the reorientation of the masticatory musculature. It is very individual how long this lasts. However, this does not have any consequences according to several studies.

<u>Periodontic pain</u>, pain in the periodontium, the splint is too tight.

Excessive salivation, the more voluminous the construction is the stronger the salivation is. After a certain time of wearing, however, a reduction of the salivation can be assumed.



Findings

The questionnaire of the Silensor-sl flyer shall help to clarify the intensity of snoring and if there might be a sleep apnea.

The result gives recommendations for a further clarification and treatment.







impairment of your health. The Silensor-sl can be fabricated

in the normal bite situation.

lower jaw in an advanced position.

Upon suspicion of a sleep apnea a specialist in sleep medicine has to be additionally consulted.



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Case example

Patient with an AHI of 76!

Almost complete reduction of the rhonchopathy with the Silensor-sl but insufficient reduction of the apneas!





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Literature/studies

Many studies with protrusion splints inclusive the Silensor-sl are available.

Statistical, many studies summarizing theses show an effectiveness against snoring of 82 % and an average reduction of the obstructive sleep apnea index of 54 %.

Treatment of snoring and obstructive sleep apnea with a mandibular protruding device: an open - label study Anette M.C. Fransson et al., sleep and breathing, vol. 5/1 2001

Oral appliances for snoring and obstructive sleep apnea: a review Kathleen A. Ferguson et al., sleep, Vol. 29/2, 2006 European Journal of Orthodontics 24 (2002) 239-249

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Mandibular advancement splints and continuous positive

airway pressure in patients with obstructive sleep

apnoea: a randomized cross-over trial

Patients N

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SUMMARY This prospective, randomized, cross-over trial was designed to compare the efficacy of a mandibular advancement splint (MAS) with that of nasal continuous positive airway pressure (nCPAP) in patients with obstructive sleep apnoea (OSA). Twenty-four patients (20 males and four females) with mild to moderate OSA (AHI between 10 and 49 events per hour) were enrolled in the study. Each patient used both MAS and nCPAP, with the initial therapy being allocated at random. Treatment periods lasted for two months with a twoweek wash-out interval between. Polysomnography was performed prior to the study and

Efficacy of Mandibular Advancement Splint for Treatment of OSA, Report at Three Months of a One-Year Follow-Up Study

Tea Galić, 🕅 talija I ković, Renata Pecotić, Joško Bžić, Tina Tičinović Kurirr Gugo Gnijača, Maja Valić, Goran Račić, Zoran Đogaš inimal jaw opening. University of Split, School of Medicine, Split, Croatia

INTRODUCTION Mandibular advancement splint Silensor-sl (MAS) can effectively treat mild to moderate obstructive sleep apnea (OSA). It is worn during sleep to maintain the patency of the upper airway by increasing its dimensions and reducing its collapsibility. Although less efficacious than continous positive airway pressure (CPAP) for improving the polysom nographic indexes of OSA, MAS is generally preferred by pati-

ents which ensures better compliance and may provide an equivalent health outcom e. MAS have been shown to have a beneficial impact on numerous dinical outcomes, including the polysomnographic indexes of OSA, subjective and objective measures of sleepiness, blood pressure, aspects of neurophysiological functioning, and guality of life. In this study we sought to evidence the efficacy of specific mandibular

advancement splint Silensor-sl and the long-term impact on numerous dinical outcomes

METHODS

- 7 patients with mild to moderate OSA
- patients were initially screened for dental status; inclusion criteria was at least 6 healthy teeth in each dental arch
- . dental impressions and lateral cephalometric radiographs were obtained prior to the initiation of the treatment arterial stiffness, blood pressure and metabolic blood parameters
- were measured at baseline and after 3 months of MAS treatment treatment outcome was determined by polysomnography





Values are given as No.(%) or mean±SD, unless otherwise indicated, BMI=body mass ndex; ESS=Epworth Sleepiness Score; STOP=snoring, tiredness, observed apnea. and hoh b ood pe ssure

RESULTS

Variables	MASt eatment	MASt eatment	P values
ESS so re	6.29±3.40	6.0±4.30	NS
AHI (events/hr)	21.79±5.78	10.76±3.98	0.0158
Minimum SpO ₂	84.0±5.35	88.57±2 p 4	NS
Mean SO 2	94.29±1.98	95.14±1.21	NS
Snoring time (min)	284.9±199.48	165.29±182.79	NS
Fibrinogen (g/L)	3.13±0.73	3.46±0.89	NS
Total cholesterol (mmol/L)	5.74±1.08	5.94±1.33	NS
Cortisol (nmol/L)	372.57±83.98	353.37±107.71	NS
FPG (mmol/L)	5.0±0.33	4.86±0.4	NS
FPI (pmol/L)	75.48±80.53	72.27±74.65	NS
HR (beats/min)	68.67±19.82	64.43±12.35	NS
SystolR: B (mmHg)	129.57±20.53	128.43±12.08	NS
Diastolic BP (mmHg)	78.0±11.58	77.14±5.37	NS
TABLE 2 EFFECTS OF		CEMENT SPUNTO	EP, RES-

PIRATION AND METABOLISM

Proton and ac induction Values are given as mean \$20 or No.(%), unless oftenvise indicated, NS-not signifi-cant; MAS-mandbular advancement splint; ESS+Eparah Seepiness Score; AH-rapneshypopnes index; SpO;-pulse oxymeter oxygen saturation; FPG-fasting plashargu.co.edi FPH-fasting pa smalt sulin; IRR-heart rate; IB-Nod pressure. P value<8.05 was c nsidered to be sit i stically is gnificant

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FIGURE 1 Whole no ht polysomnopraphy data for ce a stient/ Alice 5 dignost i c sleep sstem device; bilips R spiro A - Prior to MAS treatment B - At 3 on the 6 MAS treatmen

CONCLUSION

Mandibular advancement splint Silensor-sl may be offered as an alternative sleep. This orientatreatment with moderate improvement of OSA symptoms in patients with mild nitted only forward to moderate OSA. The significant changes in arterial stiffness, blood pressure and metabolic e during opening,

blood parameters, did not occur in 3 months of treatment but our study wil on of the airway be continued to 1-year treatment period. ndibular opening. A

REFERENCES

- 1) Chan ASL, Lee RWW, Cistulli PA. Dental appliance treatment for obstructive sleep apnea, Chest, 2007:132:693-699,
- 2) Itzaki S, Dorchin H, Clark G, Lavie L, Lavie P, Pillar G. The effects of 1-year treatment with a Herbst mandibular advancement splint on obstructive sleep apnea, oxidative stress, and endothelial function. Chest. 2007; 131:740

3) Phillips CL, Yee B, Yang Q, Villaneuva AT, Hedner J, Berend N, Grunstein R. Effects of continous positive airway pressure treatment and withdrawal in patients with obstructive sleep apnea on arterial stiffness and central BP. Chest, 2008; 134:94-100

purpose of the study, but assessment was carried out in a random order and with all patient identification removed.

The oral appliance. A soft, one-piece MAS was selected initially, similar to that described his vacuum-formed

> cheap to construct, dible forward at the

otrusion, with no

tion approximated

ble protrusion. Four-

acetate blanks were

and two laterally

buccally (Figure 1).

ie upper and lower

rnal oral breathing

lvancement of the

taking a new jaw

treated in this way,

ned of inadequate

nd were unable to

wo-part, semi-rigid

uttlingen, Germany)

remainder of the

ower elements were

ing from the upper

gions, thus allowing

nsor was its ease of

pliance.

Figure 2 Modified Silensor appliance.

adjustment. The buccal connectors are available in four lengths and the mandible may be readily advanced by replacing the original connector by a shorter one. Since modification of the splint design could have had an effect on outcome, results for the two types of splint were compared both separately and for the group as a whole.

Nasal continuous positive airway pressure (nCPAP). nCPAP was provided using the REM Star Choice machine (Respironics Inc., Medic-Aid, West Sussex, UK) at UCLH and the Sullivan Elite machine (Resmed UK Ltd, Abingdon, UK) at RBH. A comfortable nasal mask was selected and nasal corticosteroid sprays were prescribed to relieve nasal congestion if necessary. This symptom did not require treatment during the MAS arm of the study in any individual. Correct nCPAP pressures were titrated individually.

Diagnostic polysomnography. All patients had a diagnostic polysomnography before entry into the study. At UCLH, the equipment was a Compumedics system (Compumedics Ltd, Victoria, Australia), which recorded sleep and its stages by electroencephalographic (EEG), electro-oculographic, and electromyographic (EMG) criteria. EEG was recorded with electrodes placed at C3-A2 and C4-A1 (according to the international 10-20 system). EMG activity was recorded from the submental muscles. A single noir of alastropardioprophia short loads was used



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Thermoformina

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Fabrication

sl-parts card for the fabrication of a Silensor-sl.

The Silensor-sl can be fabricated with and without registration (constructive checkbite).

The sl-protrusion gauge allows the fabrication of a Silensor-sl that exactly corresponds to the measured proportions in the mouth.





The adjustment of the Silensor-sl

Select connector length: The connectors are easily exchangeable, for ex. if more protrusion is needed for a sufficient effect.

The shorter the connector is selected in relation to the measurement, the greater is the advancement of the lower jaw.



22

23

24

25

26



without bite-taking select 22 mm connector

with bite-taking select 23 mm connector

25 mm measured

without bite-taking select 24 mm connector

with bite-taking select 25 mm connector

The 26 mm connector is used when the patient despite bite-taking does not accept the advancement.



sl-protrusion gauge

The sl-protrusion gauge allows to register in a simple way the desired or recommended advancement for the Silensor-sl.



1. Marking of the normal bite situation with the sl-protrusion gauge.



2. Marking of the maximum advancement.



3. Marked desired protrusion.

Generally, half of the maximum protrusion is recommended as the advanced position of the lower jaw.



4. Registration.



Sleep disorders and dentistry

Thank you very much for your attention

An overview of the manufacturing follows on the next pages.

Fabrication (Extract of the instructions)

Model preparation

Fabrication with construction bite



In case of a very retentive teeth situation, the marking of the prosthetic equator is recommended (1.).

With the exception of the fixation points, the splint ends in case of large undercuts at the equator, otherwise 1-2 mm below.

In case of using Erkodur (hard), relieve tension from the four upper front teeth by applying Erkoskin (2.).

Block out undercuts and spaces with Erkogum, block out bubbles in the plaster with high-fusing wax. Relieve tension from the gingival margin in the area where the splint possibly has contact (3.).

If the measuring point is located on an edentulous area, this must be filled with plaster (4.). In case of a free-end situation, a plaster wall is put on the ridge (5.).



1. Separate the measuring templates.

25 mm or 23 mm ?, 4 + 5.



2. Articulate the models using a rubber band and the construction bite that has been taken off the sl-protrusee hints, page 1, paragraph sion gauge and cut to shape.



the measurement (see hints).

that a parallel drilling is possible.

occlusal plane with Erkogum. Initial point is the upper

canine or canine area. The lower pivot point results from

Fix the measuring template with the drilling shells that way

3. Fix measuring template as near as possible to the



4. Cut the spacer holding pins.

Put the marked end in the drilled holes, see 7.

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Fabrication with construction bite



5. Drill with the 1.4 mm drill (10 000 rev./min.!) through the drilling shell into the model, first in the canine area (3 mm depth of drilled hole or more).

9. Push a spacer holder as illustrated onto the spacer holding pin and press it on as near as possible. The small side always points with a knife. towards the occlusal plane.



second hole.

templates ...

10. Pay attention to a paral-

er holding pin through the

Drill the other side in the

same way. Remove pins and

Remove excessive Erkogum



7. ... the models can now be separated. Now put all 4 drill guide. Only now drill the spacer holding pins into the drilling holes. Strongly diverging spacer holding pins have to be adjusted.

Erkogum. Fix chipped plaster pieces and the pin with quick-acting glue.



8. Put a poor quantity of

Erkogum violet onto the pins.

Cut the spacers without overhang.



lelism of the modelling pads. spacer holding pin and

11. Undercuts between model have to be filled up. Now mark the outermost



surface of each spacer with

an appropriate pen.

12. Marked areas have to be 13. Articulate the models free of Erkogum. Cut all pins. with the construction bite (Erkoform-3/3d/3d motion with Occluform-3). Leave the area below the spacer at least 6 mm free of granules.





Fabrication with construction bite



14. Keep the construction bite. Lower the bite at the supporting pin to a gap of app. 2 mm between the front and close the Occluform. teeth. Pull off the insulating foil of the Erkolen foil (1.0 mm) and keep it.



15. Now thermoform, immediately put the Erkolen foil (reusable) onto the model The result is a plane occlusal surface.



16. Now take a silicone key for the opposing bite (Aton-Lab 80). Put the modelling silicone in the unit onto the splint and imprint the opposing bite with the Occluform, if necessary, slightly adapt.



of the Occluform model pot

and roughly cut out (fissure

bur > 20 000 rev./min.)



18. Lock the lower joint of the Occluform with the swivel screw. Fix lower jaw model in the Occluform model pot, fill up with granules and put the silicone key onto it. (Instructions Occluform)



19. Fix upper jaw model on the Occluform model plate. Articulate the models using the silicone key. Open the Occluform and remove the silicone key.



20. Press the cut insulating foil of the Erkolen foil with the adhesive side down on the occlusal surface of the splint.



10 mm around the spacers are free of granules.



21. Pay attention that at least 22. Now execute the second thermoforming process. As soon as the foil is adapted, close the Occluform. Allow to cool completely. Uncover all spacers before taking the splints off the model. Thereto, carefully grind through the plate ...



23. ... until the coloured

marking on the spacers is just abraded, not more and not less (tungsten carbide bur > 20 000 rev./min.). Ensure a plane surface. Take the splints off the models.

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continue at finishing

Fabrication without construction bite



1. Separate the measuring templates.

25 mm or 23 mm ?, see hints, page 1, paragraph 4 + 5.



2. Articulate the models using a rubber band.





Fix the measuring template with the drilling shells that way that a parallel drilling is possible.



4. Cut off the spacer hold-ing pins.

Put the marked end in the drilled holes, see **7**.



5. Drill with the 1.4 mm drill (10 000 rev./min.!) through the drilling shell into the model, first in the canine area (3 mm depth of drilled hole or more).



6. Immediately insert a spacer holding pin through the drill guide. Only now drill the second hole.
7. ... the models can now be separated. Now put all 4 spacer holding pins into the drilled holes.

Drill the other side in the same way. Remove pins and templates ...



7. ... the models can now be separated. Now put all 4 spacer holding pins into the drilled holes.
Strongly diverging spac-

Strongly diverging spacer holding pins have to be adjusted.



quick-acting glue.

8. Press a poor quantity of Erkogum violet onto the pins.Cut the spacers without overhang.





Fabrication without construction bite



9. Push a spacer holder as illustrated onto the spacer holding pin and press it on as near as possible. The small side always points with a knife. to the occlusal plane.

10. Pay attention to a parallelism of the modelling pads. spacer holding pin and

Remove excessive Erkogum



11. Undercuts between model have to be filled up.



12. Marked areas have to be free of Erkogum. Now mark the outermost surface of each spacer with an appropriate pen.



13. Embed the models into the granules, leave the area below the spacer at least 6 mm free of granules. Thermoform the models one after the other.



14. Immediately after the adaptation apply the Erkolen foil (1 mm) without insulating foil and press it on along the teeth row especially in the area of the front teeth, in sal surface. doing so run with the ...



15. ... finger back and forth. Do not stay too long at one place, hot!

The result is a plane occlu-



16. Take the models out of **17.** Uncover all spacers the unit and roughly cut out before taking the splints off before removing the splint from the model (fissure bur grind through the plate until > 20 000 rev./min.). the coloured marking on the spacers is just abraded, ...



18. ... not more and not less! (tungsten carbide bur the model. Thereto, **carefully** > 20 000 rev./min.) Ensure a plane surface. Take the splints off the models.



continue at finishing

Finishing



HSS twist drill (> 20 000 rev./min., without pressure), leave sufficient material (min. 2 mm) around the fixation points.



1. Cut the final form with the **2.** Grind the borders with the tungsten carbide bur (>20 000 rev./min.).



3. Smooth the borders, grinded areas with Lisko-S, narrow zones and interdental spaces with Liskoid (both 10 000 rev./min.).

4. Polish Erkodur with polishing agent for plastics (polishing set, 110 878).



5. Erkoloc-pro can be shined with the hot air burner (177 540), thereby only work on the model and do not heat the holes for the anchors (risk of deformation).



6. Press spacers inwards out 7. Remove the of the splint (for ex. with the Lisko-S mandrel shank), it might be necessary to firmly press.



insulating/shrinkage compensation foil.



8. Cut the anchors as shown 9. ... put them into the splint on the picture.

Take the anchors at the retaining lip and ...



as replacement for the spacers.



10. Firmly press into position, if necessary, also carefully with suitable pliers.



Fabrication

11. Cut the connectors, always opposing connectors have the same length.

Choose the connector length:

The connectors are exchangeable, for example if more protrusion is necessary for a sufficient effect.

The shorter the connector is chosen in comparison to the measurement, the larger is the advancement of the lower jaw.



without construction bite: measured, 23 / 25 mm connector, 22 / 24 mm with construction bite: measured, 23 / 25 mm connector, 23 / 25 mm



*The 26 mm connector is used when the patient despite of bite taking does not tolerate the advancement.



12. Remove sharp cutting edges!



13. Hinge the connectors into the long slot and pull it into its final position.



14. Observe upper jaw canine side of the connector.





Upper jaw, obligatory run of the connectors, on the left and right.



15. Hinge the connector into the other splint.Please check correct positioning of the splint.In case of propulsion movements (feed) the ...



16. ... connector shall slide out of the anchor of the lower jaw, see picture, if not, hinge the connector aboutface.



Connected splints, obligatory run of the connectors, on the left and right.





17. Now cut the retaining lips off the anchor. Finished.

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Finishing



Correct fixation of the connector

The sliding end of the connector is always placed at the lower molar area.

The fixation as shown in the upper picture is correct.

The fixation as shown in the lower picture is wrong, it may cause pain at the gingiva during sudden protrusion movements.

Also there is a risk that the connector may jump out of the anchor, due to sideward moving when sliding on the gingiva.

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Thank you for your attention.





