

Lithotripsy for Salivary Stones (for Pennsylvania Only)

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[Instructions for Use](#)

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Related Policy

- [Extracorporeal Shock Wave Therapy \(ESWT\) for Musculoskeletal Conditions and Soft Tissue Wounds \(for Pennsylvania Only\)](#)

Application

This Medical Policy only applies to the state of Pennsylvania.

Coverage Rationale

The following are unproven and not medically necessary for treating salivary stones due to insufficient evidence of efficacy:

- Extracorporeal shock wave lithotripsy (ESWL)
- Endoscopic intracorporeal laser lithotripsy

Applicable Codes

The following list(s) of procedure and/or diagnosis codes is provided for reference purposes only and may not be all inclusive. Listing of a code in this policy does not imply that the service described by the code is a covered or non-covered health service. Benefit coverage for health services is determined by federal, state, or contractual requirements and applicable laws that may require coverage for a specific service. The inclusion of a code does not imply any right to reimbursement or guarantee claim payment. Other Policies and Guidelines may apply.

CPT Code	Description
42699	Unlisted procedure, salivary glands or ducts

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Description of Services

Sialolithiasis, the formation of salivary stones due to crystallization of minerals in saliva, can cause blockage of salivary ducts resulting in painful inflammation, especially during or after meals. Most salivary stones occur in the submandibular gland, followed by the parotid gland and infrequently in the sublingual or minor salivary glands. While smaller stones may pass on their

own, larger stones generally require medical or surgical intervention. Minimally invasive gland preserving techniques of treating salivary stones have been evolving rapidly.

Extracorporeal shock wave lithotripsy (ESWL) is a minimally invasive approach that uses high energy shock waves generated outside the body to pulverize or crush the stones inside the body. Intracorporeal lithotripsy uses a lithotripsy probe inserted into the salivary duct under endoscopic guidance, and directly reaches the stone's surface and introduces shock waves in the form of laser beams, pneumatic devices and electro-hydraulic or electrokinetic probes. The goal of both techniques is to reduce the size of the sialoliths making them more easily cleared from the salivary duct system spontaneously after sialogogue-induced salivation, or during endoscopic procedures.

Clinical Evidence

There is insufficient evidence to support the use of extracorporeal shock wave and endoscopic intracorporeal laser lithotripsy for managing salivary stones. Further research with randomized controlled studies and larger patient sample sizes is required to demonstrate safety and efficacy.

In a 2020 systematic review, Chiesa-Estomba et al. sought to analyze the current literature on the role of laser assisted lithotripsy with sialendoscopy (LAS) in the treatment of major salivary gland lithiasis and provide an evidence-based perspective on clinical outcomes. Study selection criteria were as follows: reported the results for at least 10 adult patients treated for obstructive salivary gland disease via LAS, describe the type of laser device used, gland affected, stone size, outcome and complications as well as success with salivary gland preservation. The primary outcome evaluated was symptom resolution (a secondary outcome was salivary gland preservation) in a 3 month follow up timeframe. There were 16 studies that met all of the inclusion criteria, 11 were clinical retrospective non randomized uncontrolled studies, and 5 prospective non randomized studies. The most commonly reported glands were the submandibular and parotid. Results showed a success rate for stones with a mean maximum diameter of 7.11 mm ranging from 71%-100% with a gland preservation rate of 97%. The authors concluded that this systematic review suggests LAS can be an efficient, safe and gland preserving technique when done by experienced hands for the management of midsize sialolith removal from major salivary glands. However, due to the low evidence level (mostly case series without a comparison group), additional prospective-randomized trials are needed to determine the definitive role of this technique in the management of obstructive salivary gland disorders and make stronger and more precise recommendations for use of laser technology for management of not only larger stones but also other obstructive pathology such ductal stenosis.

Guenzel et al. (2019, included in the Chiesa-Estomba reviewed above) conducted a retrospective, interventional consecutive case series to evaluate the safety and efficiency of holmium laser-assisted lithotripsy during sialendoscopy of the submandibular gland. A total of 374 consecutive submandibular gland sialendoscopy procedures were performed in 276 patients with sialolithiasis from 2008 to 2015. Holmium laser-assisted Laser lithotripsy was performed in 109 cases (64.9%). Smaller mobile concretum was removed directly either by forceps or wire basket or following marsupialization of the submandibular duct. This was the case in 88 patients (29.1%). Three patients (0.8%) required surgical removal of the submandibular gland due to early abscess. The majority of patients (n=374 procedures; 90.1%) remained symptom-free after two or more years following intervention. In the remaining procedures (n=37 procedures; 9.9%), patients reported discreet postprandial problems but did not seek medical attention. The authors concluded that Holmium laser-assisted lithotripsy is a simple, safe, and effective procedure for treating patients with sialolithiasis of the submandibular gland. This study is limited by a lack of a controlled comparator group. Further investigation with prospective randomized controlled studies is needed to determine the effectiveness of laser-assisted lithotripsy in the treatment of salivary stones.

Koch et al. (2017) conducted a retrospective case series in a tertiary referral center to assess results after treatment of difficult/complex sialolithiasis with extracorporeal shock-wave lithotripsy (ESWL) and intraductal pneumatic lithotripsy (IPL). Seventy-one ESWL procedures and 57 IPL were performed in our patients. Forty-nine stones were treated by 67 ESWL procedures and 52 IPL. ESWL converted sialoliths from sialendoscopically untreatable into sialendoscopically treatable cases in 94.7%; the treatment then was completed by a total of 52 IPL procedures. ESWL was performed before IPL (81.6%), in combination with IPL (7.9%) and after (10.5%). Complete fragmentation was achieved in 97.9%. Four stones each were treated with ESWL and IPL alone in multiple sialolithiasis. Altogether, 53 stones were treated by 57 IPL procedures. Complete fragmentation was achieved in 98.1% of the 53 stones. ESWL and IPL were the dominant treatment modalities in 84.1% of all 63 stones treated. Of all 38 patients, 92.1% became stone-free and all became symptom-free. All the glands were preserved. Multiple stones were treated in 34.2% of the patients; of these, 92.3% became stone-free. The authors concluded that patients

with difficult and complex sialolithiasis can be treated with high success rates of >90% using a multimodal, minimally invasive, and gland-preserving treatment approach. ESWL and IPL played a key role in this multimodal treatment regime in > 80% of stones. This study is limited by lack of a control group and small study population.

In Capaccio et al. (2017), the authors evaluated the results of a long-term experience in the management of paediatric obstructive salivary disorders. The study involved a consecutive series of 66 children whose obstructive salivary symptoms were caused by juvenile recurrent parotitis (JRP), stones, ranula and ductal stenosis. 45 patients underwent interventional sialendoscopy for JRP, stones and stenoses, with 12 receiving a cycle of extracorporeal shockwave lithotripsy (ESWL). Other procedures included: three sialendoscopy-assisted transoral surgeries, one drainage, six marsupializations, and two suturing of a ranula. Three children underwent combined ESWL and interventional sialendoscopy, and seven a secondary procedure. Three children underwent combined ESWL and interventional sialendoscopy. An overall successful result was obtained in 90.9% of cases. None of the patients underwent traditional invasive sialadenectomy notwithstanding persistence of mild obstructive symptoms in six patients. No major complications were observed. According to the authors, using a diagnostic work-up based on colour Doppler ultrasound, Magnetic Resonance sialography and cone beam 3D TC, children with obstructive salivary disorders can be effectively treated in a modern minimally-invasive manner by extracorporeal and intracorporeal lithotripsy, interventional sialendoscopy and sialendoscopy-assisted transoral surgery; the state that this approach guarantees a successful result in most patients, thus avoiding the need for invasive sialadenectomy while functionally preserving the gland. This study is limited by the small number of the children receiving ESWL, observational design, and lack of adjustment for confounding factors.

Phillips and Withrow (2014, included in the Chiesa-Estomba reviewed above) compared outcomes and complication rates of sialolithiasis treated with intracorporeal holmium laser lithotripsy in conjunction with salivary endoscopy with those treated with simple basket retrieval or a combined endoscopic/open procedure in a cohort study. Thirty-one patients were treated for sialolithiasis. Sialoliths averaged 5.9 mm in size and were comparable between both groups. Sixty-eight percent were in the submandibular gland (n=21), with the remaining 32% in the parotid gland (n=10). Fifty-two percent of patients (n=16) were treated endoscopically with intracorporeal holmium laser lithotripsy, while the remaining 48% (n=15) were treated with salivary endoscopy techniques other than laser lithotripsy. Successful stone removal without additional maneuvers occurred in 81% of the laser cases and 93% of the non-laser group. Patients in the laser group reported an average improvement of symptoms of 95% compared with 90% of the non-laser group when adjusted for outliers. Complications in all patients included ductal stenosis (n=2) and salivary fistula (n=1). According to the authors, the results of this study show favorable outcomes with the use of intracorporeal holmium laser lithotripsy for the endoscopic management of sialolithiasis with minimal adverse events. This study was non-randomized and had a small sample size.

Desmots et al. (2014) evaluated the predictive value of sonographic fragmentation in the treatment of sialolithiasis in a case series. The main objective was to streamline the management by treating the patients with three sessions of ultrasonic lithotripsy, and to compare the success rate and complications with data from the literature. A second objective was to analyze the predictive value of data from the post procedure and follow-up sonography related to therapeutic success with regard to size, site and location of stones. The study methods included a prospective follow-up of 25 patients over a period of 31 months with one or more salivary calculi (19 parotid, submandibular 6) treated with extracorporeal lithotripsy (electromagnetic MINILITH SL 1, Storz Medical, Switzerland). No anaesthesia or analgesia was used. Each session of lithotripsy lasted on average 30 min. Complete success (absence of clinical symptoms 3 months after the end of treatment (or the last session) and residual stones <2 mm) was observed in 36% of patients, partial success (persistence of symptoms least 3 months (lower intensity and lower frequency) or size of residual stones >2 mm) in 48% and failure (persistence of same or increased symptoms at 3 months or no change in size of the calculi) in 17% of patients. Sonographic fragmentation of the stone, total energy delivered, and the total number of shock waves are predictive factors of complete success. Size, salivary topography, ductal topography, mobilization of the stones, occurrence of minor side effects and total duration of treatment had no predictive value of complete success. There was no significant difference between the first 5 and the last 20 patients. In agreement with the literature data, the efficacy of treatment was greater for parotid than submandibular calculi. The authors concluded that extracorporeal lithotripsy is an alternative to conventional surgery with no major complications. Sonographic fragmentation of calculi, total energy and total number of shock waves are predictive factors of successful treatment. This study is limited by lack of a control group and small study population.

Zenk et al. (2012) conducted a case series with chart review of 1154 patients with sialolithiasis. Diagnostic sialendoscopy confirmed 221 parotid stones and 812 submandibular stones, of which 206 and 736, respectively, were treated. Transoral stone removal was the most frequently used method to remove submandibular stones (92%). Parotid stones were removed by salivary

gland endoscopy (SGE) (22%), combined SGE and incisional technique (26%), or extracorporeal shockwave lithotripsy (ESWL) (52%), with long-term success rates of 98%, 89%, and 79%, respectively. The authors concluded that salivary gland endoscopy is an important diagnostic and therapeutic tool in the management of sialolithiasis but must be combined with additional techniques to ensure a high rate of stone clearance, symptom resolution, and gland preservation. Study limitations included no randomization or blinding and a lack of comparison with a controlled group. There was no diagnostic reference standard so a comparison between different removal methods was not possible.

In a prospective caseseries, Escudier et al. (2010) identified the factors that affect outcome (stone clearance, partial clearance without symptoms, and residual stone with symptoms unchanged) of extracorporeal shock wave lithotripsy (ESWL). The study included 142 salivary calculi (78 submandibular, 64 parotid). The results were analyzed, and a predictive model generated, which was validated using a second group of patients treated by the same technique. ESWL achieved complete success (stone and symptom free) in 67 (47.15%) of cases (submandibular 28/78, 35.9%; parotid 39/64, 60.9%). Partial success (residual stone and symptom free) was obtained in a further 49 (34.5%) (submandibular 29/78, 37.2%; parotid 20/64, 31.3%). Failure occurred in 26 (18.3%) of cases (submandibular 21/78, 26.9%; parotid 5/64, 7.8%). The investigators concluded that ESWL can eradicate salivary calculi, but its effectiveness is dependant mainly on size of the stone. This study is limited by lack of comparison group or long-term follow-up.

Nahlieli et al. (2010) assessed a combined external lithotripsy-sialoendoscopy method developed for advanced salivary gland sialolithiasis in a case series of 94 patients (43 males and 51 females). Of these 94 patients, 60 had pathologic features in the submandibular gland and 34 in the parotid gland. A miniature external lithotripter was used, combined with multifunctional sialoendoscopes and endoscopic-assisted techniques, to achieve effective removal/elimination of the stones. Total elimination of the stone using lithotripsy alone was achieved in 32% of the cases; in 29%, intraductal endoscopic assistance was needed. In the remaining 39%, the removal of a stone was achieved with the help of an endoscopy-assisted extra-ductal approach (37 cases). At 6 months of follow-up, all patients who had undergone lithotripsy or lithotripsy plus intraductal endoscopy had an absence of symptoms. Of the 37 patients who had undergone an endoscopy-assisted extra-ductal approach, 35 (95%) remained asymptomatic. The investigators concluded that lithotripsy plus intraductal or extra-ductal endoscopic treatment of sialolithiasis is a highly effective surgical method of eliminating/removing salivary stones, especially those attached to the surrounding tissue and in the secondary ducts. They also concluded that this method helps to avoid resection of the salivary glands and represents an additional development of minimal invasive surgical techniques. This study is limited by lack of a control group and lack of long-term follow-up.

U.S. Food and Drug Administration (FDA)

This section is to be used for informational purposes only. FDA approval alone is not a basis for coverage.

The FDA has approved several lithotripter devices. See the following website for information and approved indications [use product code FFK or GEX (for laser powered devices)]: <https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfPMN/pmn.cfm>. (Accessed September 8, 2020)

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Policy History/Revision Information

Date	Summary of Changes
08/01/2021	<ul style="list-style-type: none"> Created state-specific policy version for Pennsylvania (no change to guidelines)
05/01/2021	<p>Template Update</p> <ul style="list-style-type: none"> Removed <i>CMS</i> section Replaced reference to “MCG™ Care Guidelines” with “InterQual® criteria” in <i>Instructions for Use</i> <p>Application</p> <ul style="list-style-type: none"> Added language to indicate this policy does not apply to the state of Indiana; refer to the state-specific policy version
02/01/2021	<p>Application</p> <ul style="list-style-type: none"> Reformatted content Added language to indicate this policy does not apply to the state of Kentucky; refer to the state-specific policy version
11/01/2020	<p>Template Update</p> <ul style="list-style-type: none"> Reformatted policy; transferred content to new template <p>Application</p> <ul style="list-style-type: none"> Added language to indicate this policy does not apply to the state of Nebraska; refer to the state-specific policy version <p>Supporting Information</p> <ul style="list-style-type: none"> Updated <i>Clinical Evidence</i>, <i>CMS</i>, and <i>References</i> sections to reflect the most current information Archived previous policy version CS070.H

Instructions for Use

This Medical Policy provides assistance in interpreting UnitedHealthcare standard benefit plans. When deciding coverage, the federal, state or contractual requirements for benefit plan coverage must be referenced as the terms of the federal, state or contractual requirements for benefit plan coverage may differ from the standard benefit plan. In the event of a conflict, the federal, state or contractual requirements for benefit plan coverage govern. Before using this policy, please check the federal, state or contractual requirements for benefit plan coverage. UnitedHealthcare reserves the right to modify its Policies and Guidelines as necessary. This Medical Policy is provided for informational purposes. It does not constitute medical advice.

UnitedHealthcare may also use tools developed by third parties, such as the InterQual® criteria, to assist us in administering health benefits. The UnitedHealthcare Medical Policies are intended to be used in connection with the independent professional medical judgment of a qualified health care provider and do not constitute the practice of medicine or medical advice.