

Urethral Bulking System

A Minimally Invasive Treatment For Stress Urinary Incontinence





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This brochure provides information about the Bulkamid Urethral Bulking System for the treatment of stress urinary incontinence. Please read this entire brochure and discuss it with your doctor.





Glossary

Adverse event

Complication or side effect that may result from a procedure or the device

Bladder

Balloon-like organ in the lower abdomen where urine is stored

Bulkamid

Bulkamid is an injectable soft-tissue urethral bulking agent

Catheter

A temporary flexible tube to drain urine from the bladder

Bulkamid clinical study

Research study of women with stress urinary incontinence and treated with Bulkamid or another bulking agent

Continence

The term continence is used when the individual has control of their bladder

Contraindication

A medical condition that indicates Bulkamid should not be used as it may cause harm

Cystoscope

A small instrument with a viewing camera used by the doctor to view the urinary organs (urethra and bladder)

Pelvic floor muscles

The layer of muscles that support the pelvic organs and span the bottom of the pelvis

Polyacrylamide

A highly water-absorbent gel. It is the primary component of Bulkamid

Precaution

A statement in the product information that alerts the doctor to take measures to avoid a problem

Stress urinary incontinence

The involuntary leakage of urine during physical activity or exertion such as laughing, coughing or jumping

Top-up injection

When an additional Bulkamid injection is required shortly after the initial procedure to improve patient satisfaction / continence

Urethra

The tube which transports urine from the bladder to the outside of the body

Urethral bulking

The injection of material (bulking agent) into the tissues surrounding the urethra to help the urethra close to avoid accidental urine leakage. Urethral bulking does not completely close the urethra; it can still open normally to allow for urination

Urge urinary incontinence

Urge incontinence happens when you have a strong need to urinate but can't reach the toilet in time

Urinary incontinence

The unwanted and involuntary leakage of urine

What is stress urinary incontinence?

Stress urinary incontinence is the unintentional passing of urine during activity or exertion, such as during coughing, laughing, or exercise. It is caused by a weakness of the *pelvic floor muscles*.

Stress urinary incontinence is a common condition that affects over 13 million adults in the United States, 85% of which are women.¹ This condition can have a significant impact on daily life, affecting activities, relationships and emotional well-being. It can occur at any stage of life, with pelvic disorders from childbirth, pelvic surgery and aging increasing the risk of incontinence.

How is stress urinary incontinence normally treated?

Stress urinary incontinence is a very treatable condition. Your doctor will advise you of the different options available and help you make the right treatment decision. These may include:

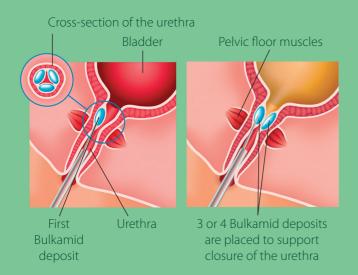
Pelvic Muscle Strengthening

Pelvic floor exercises, commonly referred to as Kegel exercises, will most likely be one of the first treatment options recommended by your doctor. These exercises help improve support of the *bladder* and *urethra*.

Bulking Agents

Urethral bulking is the injection of a bulking agent, like *Bulkamid*, into the *urethra* to add volume ("bulk") to the tissue. It is less invasive than surgery. A bulking agent supports the closing mechanism of the *urethra* and provides better control of urine when you cough, laugh, exercise or change position.

Urethral bulking does not close the *urethra* totally; the *urethra* still opens normally to allow for urination.



Surgery

Surgery is often used to treat *stress urinary incontinence*, especially when the cause is a weakened pelvic floor. A common surgery to provide support for the *urethra* is a sling or mesh procedure. During this surgery, a narrow strip of a permanent material is placed below the *urethra* to support it and to maintain its seal.

All treatment options have *contraindications, precautions* and *adverse events* (side effects) associated with their use. You should discuss the different treatment options available with your doctor.

What is Bulkamid?

Bulkamid is a **urethral bulking** agent, consisting of 97.5% water and 2.5% **polyacrylamide**. **Bulkamid** is injected into the soft tissue of your **urethra** using a syringe. **Bulkamid** achieves its bulking effect by the volume of the gel injected; other bulking agents achieve their bulking effect by a combination of the micro particles themselves and the body's own reaction to the particles.

Bulkamid is indicated for the treatment of women who have *stress urinary incontinence (SUI)* or *stress predominant mixed incontinence* (combination of SUI and urge incontinence).

When should Bulkamid not be used?

Bulkamid Urethral Bulking System must not be used in patients suffering from acute urinary tract infection.

Bulkamid should not be used in male patients.

For additional information about safety and risks, please see page 13 or go to www.bulkamid.com/ISI

What were the results of the Bulkamid Clinical Study?

A *Bulkamid clinical study* was conducted at 33 medical centers in the United States and Canada. Two hundred and twenty eight (228) women with *stress urinary incontinence* received *Bulkamid* and were followed for 12 months after the initial treatment; 46.9% (107 of 228) of these women reported their symptoms to have improved by at least half (50%), while three quarters (75%) of women reported dryness or improvement in their symptoms after *Bulkamid* treatment.²

In the *clinical study*, once women had received the initial injection of *Bulkamid*, they were offered up to two additional injections if their symptoms were still present. Seventy-seven (77) percent (176 of 228) of women treated with *Bulkamid* had additional injections of *Bulkamid*.²

The most common side effects (*adverse events*) reported by *Bulkamid clinical study* patients are listed in the table below. Most of these *adverse events* were temporary. One patient treated with *Bulkamid* experienced a serious *adverse event* (blood in urine) that was thought to be related to the bulking procedure but not to the *Bulkamid* gel.



Most common side effects reported in the Bulkamid Clinical Study

| Side Effect | Patients |
|--|-----------|
| Pain at implantation | 13 in 100 |
| Acute urinary retention | 6 in 100 |
| Urinary tract/ bladder infection | 4 in 100 |
| Blood in urine (hematuria/ transient hematuria) | 2 in 100 |
| Painful urination (dysuria) | 1 in 100 |
| Strong desire to urinate, but no incontinence episodes (urgency) | 1 in 100 |
| Urge incontinence | 1 in 100 |
| Worsening <i>urinary</i> <i>incontinence</i> | 1 in 100 |
| Vaginal infection | 1 in 100 |

What are the risks of the Bulkamid procedure?

Over 70,000 women with *stress urinary incontinence* in 25 countries over 10 years have been treated with *Bulkamid*. During that time, a low number of complications have been reported and there have been no reported long-term complications. However, as with any invasive procedure, complications may occur.

Complications can include pain related to the procedure, a small amount of blood in the urine, delayed urination, painful urination, and/or urinary tract infection. These complications are usually temporary and normally resolve within a few days. In very rare cases patients may experience difficulties to normally pass urine and may require the use of a disposable catheter to catheterize themselves to empty the bladder a few times each day until the bladder starts to empty on its own again.

Although not seen in the *Bulkamid clinical study*, the following side effects have occurred in women who received other urethral bulking agents: blood vessel blockage (embolism/vascular occlusion), skin redness (erythema), loss of bulking agent material through the injection site (excreted material), movement of bulking agent material to a different location inside the body (migration), inflammation causing a growth of tissue (granuloma), inability to completely empty the bladder (outlet obstruction), loss of urethral tissue over the implant site (erosion), and frequent urination.

There is a possibility that you may not experience any benefit from *Bulkamid* treatment. This could happen for various reasons, such as *Bulkamid* being placed too deeply in the tissue, thereby creating poor bulking around the *urethra*. Also, if you have a different type of incontinence, such as *urge incontinence* or if your incontinence condition worsens, *Bulkamid* may not be an effective treatment for you.

What are the benefits of the Bulkamid procedure?

The majority of women treated with *Bulkamid* report dryness or improvement in their symptoms, with many seeing that improvement as soon as they leave the doctor's office, hospital or clinic. While being free from unwanted leakage is the most desired outcome of treatment, many women consider a successful treatment to be a decrease in the amount and frequency of urine leakage due to *stress urinary incontinence.*

If relief from your symptoms is not sufficient following treatment with *Bulkamid*, an additional injection (a *"top-up" injection*) can be given to help achieve satisfactory results. It is recommended that you wait at least 4 weeks between treatments to realize the full benefit of the first treatment. Talk to your doctor about an additional treatment if you continue to experience urine leakage after the first treatment.

The benefit of Bulkamid treatment is that you could be free from unwanted urinary leakage or have fewer episodes of urinary leakage. In the *Bulkamid clinical study* women were asked how effective they felt their treatment was 12 months after their initial injection. Over three quarters of women reported that their incontinence was either "dry, much improved or improved" and approximately half of women reported zero stress urinary incontinence episodes.²

What can I expect on the day of my treatment?

The *Bulkamid* procedure is minimally invasive and typically takes about 10-15 minutes to perform. The procedure usually takes place in an outpatient clinic and you will normally be able to go home on the same day.

Prior to the procedure, your doctor will discuss whether you should have a local or general anesthetic to reduce any discomfort associated with the procedure. Most patients will undergo a *Bulkamid* procedure under local anesthetic and will feel no more than a slight scratch as the needle enters the urethral wall. Your doctor may prescribe an antibiotic to reduce the risk of infection.

During the procedure a *cystoscope* (a small instrument with a viewing camera used by the doctor) will be inserted into the *urethra*, which allows the procedure to be completed under constant visualization. Three or four deposits of *Bulkamid* (1.5 – 2 mL total volume, equivalent to slightly less than half a teaspoon) will be placed into the urethral tissue until the opposite walls are gently touching in the center of the urethral opening. The *cystoscope* is removed after the injection is complete and your treatment is finished.

What can I expect after the procedure?

After the procedure, your ability to empty your **bladder** will be checked prior to leaving the clinic to ensure no further assistance is required.

If you cannot pass urine, a *catheter* may be required for a short period (normally less than 24 hours but in very rare cases it may take days to weeks) to allow emptying of the *bladder*. This is an uncommon complication occurring in only 6 out of every 100 patients. Once at home, patients return to normal activities within 24 hours, depending on their doctor's advice.

If you need to undergo a surgical procedure in the future, especially a dental procedure, there is a risk of infection developing near where the *Bulkamid*

was placed. This is due to the possibility of bacteria moving and settling into the implant, which is the case with many other medical implants. So if you require a surgical or dental procedure in the future, please tell the treating physician that you have a permanent implant and discuss the need for possible antibiotic therapy with them.

For more information about Bulkamid, please visit www.bulkamid.com

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In the event of an incident with Bulkamid contact customersupport@axonics.com

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