

**Aesthetic Surgery Journal - 2002**

After successfully treating breast implant capsular contracture with ultrasound, the author asks, "If demonstrated that ultrasound is effective for treating already existing contractures, could it be also effective preventing them?" Here he presents his protocol and preliminary results of prophylactic application of ultrasound for the avoidance of capsular contractures. (Aesthetic SurgJ 2002;22:205-207.)



**Jorge Planas,  
MD, Barcelona,  
Spain, is a plastic  
surgeon**

The causes of breast implant capsular contracture are unclear and most likely multifactorial.<sup>1-3</sup>

Although implantation of textured surface implants<sup>4-7</sup> and several drug administration regimens<sup>8-13</sup> diminished the percentage of contractures, they still occur. Six years ago, I started applying external ultrasound to treat breast implant capsular contractures. Preliminary results were so positive that I was encouraged to continue.<sup>14,15</sup>

The ultrasonic device that I use is similar to the one used for superficial soft tissue treatment. In an early study I analyzed 52 patients, 25 of whom had bilateral contractures. Nineteen percent of the implanted breasts had grade IV Baker scale contracture, whereas the remaining 81 % were distributed between Baker scale grades III and II. The number of treatment sessions was determined by evaluating improvement. Patients were treated with repeat ultrasonic applications, ranging from 2 to 16 sessions, with an average of 6.4 sessions.<sup>15</sup>

To measure the effect of external ultrasound, contracture grade was analyzed before and after treatment. Changes were measured by subtracting the Baker scale value of the final state from the initial one. In all cases, a positive difference indicated an improvement in the patient's condition. In this study, I obtained an overall improvement rate of 82.6% at 1-year follow-up, with almost half of the contractures reaching total softness (Table 1).

Measurement	Pretreatment distribution %	Post-treatment distribution %
Baker I		48
Baker II	34	40
Baker III	47	8
Baker IV	19	4

table 1. Measurement of capsular contractures around breast implants. Pretreatment Post-treatment

In a preceding study of 24 Patients, 14 treated similarly, I found that in 97% of cases the degree of contracture improved at least 1 Baker degree. Joining both studies, an evaluation of 83.8% improvement at 1-year follow up confirms observations of capsular softening and easier closed capsulotomy after external ultrasound treatment. In most cases, a limited number of sessions, fewer than 8, was enough to obtain a long-term result. A satisfactory result was obtained in 75% of the cases. I also confirmed that the percentage of improvement was higher in patients with prepectoral-placed implants.<sup>15</sup>

The external ultrasonic treatment has proved to be easy to apply, well accepted by patients, and free of significant complications.<sup>14,15</sup>

After analyzing the data and considering the positive results, I posed the following question: if I demonstrated that ultrasound is effective for *treatment* of already existing contractures, could it be effective in *preventing* them? Theoretical justification for prophylactic use is based on demonstrated properties and effects (Table 2).<sup>16-18</sup>

**Mechanical** Produces micromassages that improve lymphatic drainage and help to resolve the edema

**Thermal** Increases speed of cellular metabolism  
Activates fibroblast production  
Helps the healing process, arranging the scar architecture

**Biochemical** Helps vascular proliferation  
Increases tissue oxygenation  
Increases release of cellular mediators of inflammation  
Increases fibrolytic processes

Table 2. Effects of ultrasound

We theorized that early application of ultrasound facilitates healing, diminishes edema, and regulates inflammation, thereby diminishing the possibility of a future capsular contracture. The following prophylactic application was suggested initially: session 1, 24 hours after surgery; session 2, 3 days after surgery; session 3, 7 days after surgery; and session 4, 1 month after surgery.

Ultrasound was administered under the following parameters: level, prophylactic; energy, 60 J; power, 12 W; type, pulsed; time, 10 minutes. Early application of external ultrasound was associated with the highest postsurgical inflammatory peak. The later applications were administered according to a "modulation" scheme of inflammation for up to 3 months when healing and remodeling of collagen were already established.<sup>19-22</sup>

Generally, in the first session patients did not report any discomfort; however, in the second session (third postoperative day), some patients reported hypersensitivity, mainly in the submammary fold. In the following sessions, patients did not complain of discomfort.

Currently, I have modified the number of sessions and the schedule of application as follows: session 1, 7 days after surgery when removing the stitches; session 2, 15 days after surgery; session 3, 21 days after surgery.

This new protocol avoids the hypersensitivity that some patients had in early sessions by starting treatment when the capsule around the prostheses is already constituted.<sup>19-21</sup>

It has been almost a year and a half since I began using this prophylactic protocol, and the preliminary results demonstrate faster reduction of edema and inflammation, faster absorption of small bruises and ecchymoses, and a decrease of postsurgical discomfort. Most important is that from the first patients receiving this treatment to the current patients, none has experienced the formation of capsular contracture thus far.

In view of these good results, I have followed this protocol and improved its design, and I look forward to statistically validating the different variables. At the moment, I am carrying out both protocols in parallel: therapeutic and prophylactic. Therapeutic results are quite encouraging and prophylactic results fulfill our expectations so far.

## References

1. Burkhardt BR. Capsular contracture: hard breasts, soft data. *Gynecol Plast Surg* 1988;15:521-532.

2. Georgiade NO. *Aesthetic Surgery of the Breast*. Philadelphia: WB Saunders Co; 1990.
3. McCarthy JG. *Plastic Surgery*, Vol. VI. Philadelphia: WB Saunders Co; 1990.
4. Burkhardt BR, Eades E. The effect of Biocell texturing and povidone-iodine irrigation on capsular contracture around saline-inflatable breast implants. *Plast Reconstr Surg* 1995;96:1317-1325.
5. Handel N, Jensen A, Black Q. The fate of breast implants: a critical analysis of complications and outcomes. *Plast Reconstr Surg* 1995;96:1521-1533.
6. Lesesne CB. Textured-surface silicone breast implants: histology in the human. *Aesth Plast Surg* 1997;21:93-96.
7. Rioja Torrejón L, Redondo A, De No L, Benitez J. Estudio comparativo de las complicaciones de implantes texturados rellenos de gel de silicona en oposición a los de relleno de suero. *Gir P/ast Ibero-latinoamericano* 1998;4:395-401.
8. Berman B, Duncan MR. Pentoxifyline inhibits normal human dermal fibroblast in vitro proliferation, collagen, glycosaminoglycans, and fibronectin production, and increases collagenase activity. *J Am Acad Dermatol* 1989;21:605-610.
9. Berman B, Duncan MR. Pentoxifyline inhibits the proliferation of human fibroblasts derived from keloid, scleroderma, and morphea skin, and their production of collagen, glycosaminoglycans and fibronectin. *J Am Acad Dermatol* 1990;23:339.
10. Caffee HH, Rotatori DS. Intracapsular injection of triamcinolone for prevention of the contracture. *Plast Reconstr Surg* 1994;94:824-828.
11. Ellenberg AH. Marked thinning of breast skin flaps after the insertion of implants containing triamcinolone. *Plast Reconstr Surg* 1977;60:755-758.
12. Marin Bertolin S. Profilaxis de la contractura capsular mediante pentoxifilina intraprotésica: estudio experimental en ratas. *Gir P/ast Ibero-latinoamericano* 1997;4:373-381.
13. Spear SL, Matsuba H, Romm S, LiUle JW. Methyl prednisolone in double-lumen gel-saline submuscular mammary prostheses: a double-blind prospective, controlled clinical trial. *Plast Reconstr Surg* 1991;87:483-487.
14. Planas J, Migliano E, Wagenfuhr J Jr, Castillo S. External ultrasonic treatment of capsular contracture around breast implants. *Aesth Plast Surg* 1997;21:395-397.
15. Planas J, Cervelli V, Planas G. Five years experience on ultrasonic treatment of breast contractures. *Aesth Plast Surg* 2001;25:89-93.
16. Carpaneda CA. Inflammatory reaction and capsular contracture around smooth silicone implants. *Aesth Plast Surg* 1997;21:110-114.
17. Lehmann JF, De Lauter BJ. Diatermia y Terapéutica superficial con calor, láser y frío. in Krusen J, 4th ed. *Medicina física y rehabilitación*. Madrid: Editorial Medica Panamericana; 1994.
18. Scott WW, Scardino PL. A new concept in the treatment of Peronyie's disease. *South Med J* 1984;41:17-19.
19. Batra M, Bernard S, Picha G. Histologic comparison of breast implant shells with smooth, foam, and porous microstructuring in a rat model from 1 day to 6 months. *Plast Reconstr Surg* 1995;95:354-363.
20. Lilla JA, Vistnes LM. Long-term study of reactions to various silicone breast implants in rabbits. *Plast Reconstr Surg* 1976;57:637-649.
21. Vistnes LM, Kasander GA, Kosek J. Study of encapsulation of silicone rubber implants in animals. *Plast Reconstr Surg* 1978;62:580-588.
22. Wyatt LE, Sinow JD, Wollmann JS, Sami DA, Miller T. The influence of time on human breast capsular histology: smooth and textured silicone-surfaced Implants. *Plast Reconstr Surg* 1998;102:1922-1931.

