Stahl’s ear deformity is an uncommon congenital auricular anomaly that can lead to psychological disruption. It is mainly characterized by a third crus in the auricular cartilage, extending from the antihelix to the helix crus, giving rise to an irregular helical rim and a bent scaphoid fossa. This causes the top of the ear to appear pointed and flat. It is usually corrected surgically during childhood. Alternatively, the ear molding technique has been shown to be a safe and effective nonsurgical treatment. Choosing a conservative treatment over otoplasty includes fewer complications and is less stressful for both parents and children.

Nonsurgical procedures for the correction of congenital ear deformities were first published during the 1980s in Japan by plastic surgeons, and since then their use has been popularized. It is now acknowledged that keeping the ear in a constant position for a specific period will result in a normal ear shape. That happens because of the softness and lack of elasticity that the auricle of the neonate has right after birth. Physiologically this can be explained by the increased estrogen production during neonatal period, which results in elevated hyaluronic acid levels. The increased levels of the former leads to the separation of the intercellular materials (collagen, elastin, proteoglycans) from the chondrocytes, thus the cartilage lacks elasticity. During this fundamental period, when the ear is softer and more malleable, the application of constant pressure on the deformed cartilage will give the ear a more normal shape. That is why the sooner the intervention begins, the shorter the application period will be.

**CASE PRESENTATION**

A 1-week-old neonate was brought by the parents to the plastic surgery ward because of an ear deformity dating from birth, with no pathological history according to the parents’ report. The female child was born at full term. Clinical examination revealed a unilateral Stahl’s ear deformity (Fig. 1A). It was mainly characterized by a third crus and a deformed scaphoid fossa. A splinting device (EarBuddies) was applied on the same day (Fig. 1B).
The newborn was managed with a nonsurgical method by applying an ear molding device, on the 7th day postpartum. A bendable splint made of medical grade plastics, silicones, and inert metals was placed, and left in position for a period of 3 weeks. Starting just above the triangular fossa, the splint was applied to the scaphal hollow inside the rim. After that, the splint was stabilized with the use of hypoallergenic, latex-free, and water-resistant adhesive tapes, which were changed once a week or when needed. The splinted ear was further supported using a bigger tape attached to the head. The parents were instructed to readjust the tapes when displacement of the molding device was observed. During bath time, a small plastic shower cap was used to keep the area dry. Regular follow-up visits were done every 7 days. An improvement of the appearance of the ear was notable on day 12. Regarding the outcome, the affected auricle regained its normal appearance (Fig. 1C). The parents were satisfied with the result. Long-term follow-up showed no recurrence of the deformity (Fig. 1D).

**DISCUSSION**

According to a large review of 1000 infants, congenital auricular deformities are found in 55.2% of neonates at the neonate's auricle. (A) Stahl's ear before correction. (B) Molding device stabilized with adhesive tapes. (C) The affected auricle regained its normal appearance 3 weeks later. (D) Long-term follow-up at 7 months.
birth, with Stahl’s ear appearing in only 8.7% of all newborns. Stahl’s ear deformity is characterized by a misshaped but fully developed pinna, and it is only developed perinatally. Postpartum splinting can be performed in many ways, as long as the desired shape of the auricle is maintained in position with continuous pressure. A variety of techniques for surgical reconstruction of this deformity have been described, but nonoperative methods also have satisfactory results, mainly when used during the neonatal period. The complications of an otoplasty (hematoma, bleeding, and postoperative infections) are absent, and as Smith et al reported, there is a 100% correction rate for Stahl’s ear deformity. Moreover, Byrd et al showed a correction rate of more than 90%, which was significantly higher than the self-correction rate (30%). Additionally, the noninvasive method prevents later emotional distress for both parents and the child, as the deformity is treated right after birth, in a painless and inexpensive way.

Over the past few years, splinting congenital ear deformities in early neonatal life is becoming even more popular; however, it needs to be better brought to the attention of the scientific community. Firstly, it is fundamental that health care providers should be properly educated about early detection and efficient management of the ear deformity. Furthermore, guidance regarding the correct application of splinting should be given to the parents. It is crucial that neonatal pediatricians must establish a full physical examination, including detailed auricular inspection, and once the diagnosis is reached, a referral to a plastic surgeon must be suggested as soon as possible. Any delay may result in a possible unneeded surgery in the future.

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