Vacuum-assisted cesarean section

Abstract: There has been a dramatic rise in the frequency of cesarean sections, surpassing 30% of all deliveries in the US. This upsurge, coupled with a decreasing willingness to allow vaginal birth after cesarean section, has resulted in an expansion of the use of vacuum assistance to safely extract the fetal head. By avoiding the use of a delivering hand or forceps blade, the volume being delivered through the uterine incision can be decreased when the vacuum is used properly. Reducing uterine extensions with their associated complications (eg, excessive blood loss) in difficult cases is also a theoretical advantage of vacuum delivery. Maternal discomfort related to excessive fundal pressure may also be lessened. To minimize the risk of neonatal morbidity, proper cup placement over the “flexion point” remains essential to maintain vacuum integrity and reduce the chance of inadvertent detachment and uterine extensions. Based on the published literature and pragmatic clinical experience, utilization of the vacuum device is a safe and effective technique to assist delivery during cesarean section.

Keywords: cesarean section, vacuum, forceps, birth, delivery

Introduction

The rate of cesarean delivery has increased dramatically worldwide over the past several decades and now exceeds 55% of all deliveries in many countries. In the US, cesarean section frequency has surpassed 30% for nearly a decade, with a wide distribution that ranges from 7.1% to 69.9% across hospitals. Despite public health efforts to optimize and curtail cesarean section utilization, delivery rates by this method continue to rise unabated. Consequently, physicians will increasingly encounter a wider variety of clinical presentations that necessitate availability of the full range of delivery options to manage different medical scenarios and maximize outcomes.

Once established as a safe and effective method of operative vaginal delivery, vacuum-assisted procedures have more recently shown clinical utility and have become widely adopted for cesarean section. In an elective cesarean section, the lower uterine segment is commonly not elongated or effaced by labor, making it difficult to create an adequate incision to enable an uncomplicated delivery. In addition, at the time of an elective repeat cesarean section, the fetal head is normally not deeply engaged in the pelvis. Procedures to facilitate delivery in this situation include fundal pressure, forceps, internal podalic version, or addition of a lateral vertical incision (“J” incision), all of which can be traumatic for both mother and fetus. The advantages of using a vacuum device in this situation to assist delivery of the fetal head are listed in Table 1.

Historical perspective

To identify published literature relevant to vacuum-assisted cesarean section, a computer-based electronic search of the PubMed and Web of Science databases up to
and including May 2016 was conducted. The search strategy included common words and terms related to vacuum-assisted deliveries, especially during cesarean section. Also reviewed were unpublished studies to determine whether there were data sets that may be informative. Finally, bibliographies of all relevant papers were hand searched for applicable articles.

Having already achieved widespread clinical acceptance as a mainstay procedure in operative vaginal deliveries, vacuum-assisted cesarean section was first described by Solomons in his seminal report of 20 consecutive cases delivered with the 5 cm Malmström cup. All 20 infants were successfully delivered by this method, and the occurrence of caput succedaneum (chignon), typically produced by the vacuum cup, was much less apparent in these newborns. It was hypothesized that this was due to the less time and traction force required to deliver via an abdominal incision compared to the effort required to deliver the fetal head across the pelvic floor as with operative vaginal delivery.

Also using the original Malmström cup, Bercovici successfully delivered 20 of 22 (91%) term infants via cesarean section without uterine extensions. A small chignon was noted on all infants, which resolved within 2 days. In one case, delivery failure was ascribed to use of a larger (6 cm) cup and, in the other case, to a deeply engaged head that was manually delivered. It was concluded that the vacuum was most appropriately indicated for cases where the fetal head is not deeply engaged in the pelvis and does not require manual elevation out of the pelvis prior to the application of the cup.

Pelosi and Apuzzio published the initial report of the use of a soft vacuum cup (silastic) for cesarean section. Thirty-five infants were successfully delivered with this method, including 15 primary nonelective cesarean sections without evidence of fetal distress and 20 elective repeat procedures. While there were no uterine extensions or surgical complications in their series, the uniformly encouraging results could not be duplicated by others and, in fact, the popularity of the esthetically pleasing soft vacuum cups has declined substantially given the high detachment and overall failure rates.

In a prospective trial of 18 term, elective cesarean sections, Arad et al compared neonatal outcomes between eight patients delivered via vacuum and 10 patients via manual extraction. There were no notable or statistically significant differences between groups for birth weight, Apgar scores, or acid/base status measured from umbilical cord blood gases. The time required from uterine incision to complete delivery was approximately twofold higher in the vacuum group (79.4 vs 40.9 seconds, \( P<0.01 \)), but the difference was not clinically significant. Additionally, Arad et al retrospectively evaluated neonatal outcomes in 78 term, elective cesarean sections performed at their facility over a 4-year period. Again, there was no difference in Apgar scores between 34 infants delivered with vacuum assistance and 44 randomly selected infants delivered manually. These results were corroborated by Boehm, who reported successful delivery of 50 cases using vacuum-assisted cesarean section, with no neonatal morbidity.

In a randomized controlled trial, Bofill et al compared outcomes among 15 patients delivered with vacuum, 14 manually, and 15 by forceps. In the vacuum group, 13 of 15 (87%) were delivered successfully and the remaining two were delivered manually. One failure was due to a pump malfunction and the other was inadvertently manually delivered when the fetal head was being manipulated prior to cup application. In the manual delivery group, 11 of 14 (79%) were delivered successfully, while three required vacuum assistance to facilitate delivery. Delivery with the aid of forceps was successful in 11 of 15 (73%) cases; however, three had to be delivered manually and one with vacuum. There were no differences noted across groups for the Apgar scores, cord arterial pH, birth weights of the neonates, extensions of the uterine incision, or the pre- to postdelivery hematoglobin values. The time required to effect delivery tended to be longer in the forceps group \( (P=0.06) \). Importantly, patients reported less pain during delivery in the vacuum group \( (P=0.02) \) probably due to the reduced fundal pressure necessary to effect delivery.

Dimitrov et al conducted a prospective study comparing vacuum-assisted delivery \( (n=19) \) with standard manual extraction \( (n=25) \) in patients having an elective cesarean section. There were no differences in Apgar scores between groups. However, the duration of scalp traction was significantly shorter than the time necessary for manual extraction \( (30\pm4 \text{ vs } 53\pm21 \text{ seconds}, P<0.01) \). The authors noted that vacuum delivery was accomplished in all cases without the need for additional fundal pressure.

Most recently, Sritippayawan and Chantrapitak reported the results of a large randomized controlled
trial of vacuum-assisted delivery (n=90) compared to manual extraction (n=90) during elective cesarean section. Using a soft cup to assist in the delivery of the unengaged fetal head, the delivery duration was significantly shorter in the vacuum group compared to the manual delivery group (65.3±31.2 vs 86.3±53.9 seconds, P<0.01). Mean blood loss was slightly higher in the vacuum group (576.7±182.9 mL vs 504.4±204.9 mL, P=0.31), and there was no difference in Apgar scores and no evidence of chignon. The authors concluded that extraction of the fetal head by cesarean section with the vacuum device was safe, rapid, and nontraumatic and did not require the need for prolonged fundal compression.

**Current state of the art**

The original metallic Malmström cup vacuum device served as the foundational design for the current-generation rigid cup devices appropriate for use during cesarean section. One such device is the Kiwi® OmniCup® (Clinical Innovations, Salt Lake City, UT, USA) which uses a 5 cm rigid plastic cup (Figure 1). This complete vacuum-assisted fetal delivery device is designed for use in all fetal head positions—occiput anterior, occiput posterior, and occiput transverse—and cesarean section procedures. Its flexible stem and low-profile cup enable placement over the flexion point, 3 cm forward of the posterior fontanelle along the sagittal suture, so that the smallest diameters of the fetal head are presented to the birth canal or the uterine incision depending on the procedure it is being used for.

The Kiwi® OmniCup® was evaluated in a prospective trial comparing the safety and effectiveness of delivery of the fetal head during cesarean section compared to the traditional method of manual extraction (McQuivey, unpublished data). Twenty-five patients were allocated to each treatment arm. Estimated blood loss was lower in the vacuum group (681 vs 810 cc), but there were no differences in neonatal outcomes such as Apgar scores or birth weight (3.5 kg vs 3.6 kg, P=0.6), between groups. In the manual extraction group, four patients (16%) required uterine extensions and one had evidence of a cervical laceration. No uterine extensions or cervical lacerations occurred in the vacuum group. There were three reported cases of cosmetic scalp effects, such as bruising or presence of a chignon, in neonates delivered with vacuum, although all were judged minor and resolved spontaneously within 1 week. In these three instances, the cup was not positioned correctly over the flexion point and pop-offs occurred. In cases where the cup was appropriately placed over the flexion point and the integrity of the vacuum was maintained without pop-offs, there was no evidence of scalp effects.

**Discussion**

While some early adopters promoted the commonly held misconception that little skill is required to use the vacuum device, we offer a more cautious perspective given the preponderance of clinical experience. 1) Based on the formative work by Solomons and Bercovici, those infants who are unengaged or “floating” more readily lend themselves to the use of the vacuum during cesarean section. 2) Awareness of the fetal head position is of the upmost importance in allowing the surgeon the best opportunity to position the cup properly to ensure vacuum integrity without inadvertent cup detachments. In accordance with the instructions provided by Boffill et al, the cup should be placed over the flexion point, ~3 cm anterior to the posterior fontanelle along the sagittal suture. This is the point at which the mentovertical diameter crosses the sagittal suture, promoting flexion of the fetal neck and thus presenting the smallest diameter of the fetal head to the uterine incision. 3) Posterior-designed cups allow easier maneuverability of the vacuum device over the flexion point within the confines of the uterine incision. 4) The traction force should be minimized and the extent of the incision should not limit safe passage of the fetal head and body. Additional contraindications of vacuum delivery include prematurity (<36 weeks gestation), a known fetal blood or bone demineralization disorder, or known maternal viral infection. 5) Although less likely to occur during cesarean section, use of the vacuum should be discontinued after two pop-offs have occurred or after 20 minutes have elapsed.

There have been few neonatal adverse events reported in the literature related to the use of vacuum-assisted cesarean section. However, complications such as scalp abrasions,
retinal hemorrhages, jaundice, and cephalohematomas have been reported with vacuum-assisted operative vaginal delivery. More serious complications can include intracranial hemorrhages and skull fractures; however, these rare events are likely associated with misplacement of the cup, excessive traction force, and multiple pop-offs during operative vaginal deliveries. As speculated by Solomons, the force required to deliver the fetal head through the abdominal incision is less than the force required to deliver the head across the perineal floor during a vaginal delivery; thus, excessive tractions and pop-offs should be rare with cesarean section. To reemphasize, if there is difficulty delivering the fetal head through the abdominal incision, immediate attention should be paid to the sizes of the skin, fascial, and uterine incisions. The size of the incision should not be limited at the expense of safely delivering the fetus.

The combination of increasing cesarean section use and the decreasing willingness to allow vaginal birth after cesarean section will likely result in an expansion of the use of vacuum assistance. When the vacuum device is used appropriately, the delivery can be facilitated by effectively decreasing the volume delivered through the uterine incision due to the avoidance of a delivering hand or forceps blade. The vacuum may lead to decreased uterine extensions and decrease in blood loss associated with efforts to deliver the head in difficult cases. Without the need for excessive fundal pressure, maternal discomfort can be minimized. Based on the published literature and pragmatic clinical experience, the use of the vacuum device is a safe and effective technique to assist delivery during cesarean section.

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Disclosure
Dr McQuivey is an employee of and Dr Block is an independent advisor to Clinical Innovations (Salt Lake City, UT, USA). The authors report no other conflicts of interest in this work.

References