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Original Research Article

History-indicated transvaginal cerclage: results from a single-centre

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ABSTRACT

Background: Cervical incompetence occasionally results in mid-trimester pregnancy loss, preterm labour and increased foetal morbimortality. History-indicated cerclage is proposed when obstetric history suggests cervical incompetence. The aim of this study was to evaluate the maternal-foetal outcomes following prophylactic cervical cerclage.

Methods: Retrospective study reviewing data of all women undergoing transvaginal history-indicated cerclage from January 1st, 2008 to December 31th, 2017 at Centro Hospitalar Universitário do Algarve - Faro. Primary outcome: gestational age <37weeks at birth. Secondary outcomes: neonatal morbimortality and intensive care unit (NICU) admission and maternal morbidity. Data were analyzed with IBM SPSS Statistics 23.

Results: A total of 12 history-indicated cerclages were performed (9 women). At first cerclage, mean maternal age, gestity, parity and live children were 27.6, 2.44, 1.11 and 0.78 (87.7% preterm), respectively. At cerclage placement, mean gestational age and cervical length were 16.1 weeks and 27.5mm. Average hospital admission was 10.7 days. In all cases McDonald technique was performed. Four hospital readmissions occurred for threatened labour. Mean gestational age at cerclage removal was 36.9 weeks (83.3% in ambulatory) and 38.9 at delivery. Average time between cerclage removal and labour was 14.5 days. Spontaneous onset of labour occurred in 75% and vaginal delivery in 83.4%. There were no reports of preterm birth, foetal admission to NICU or maternal complications. Mean number of live children after procedure was 1.58.

Conclusions: Prophylactic cervical cerclage seems to improve pregnancy outcome with minimal maternal risks. However, our data suggest over inclusion of women, with unnecessary procedures, emphasizing the importance of re-evaluating inclusion criteria.

Keywords: Cervical cerclage, Cervical stitch, Cervical insufficiency, Cervico-isthmic incompetence, Preterm birth, Pregnancy loss

INTRODUCTION

Prematurity is a major cause of neonatal and infant morbidity and mortality.¹ Several causes, maternal and foetal, can originate cervical modifications in late second and early third trimester and result in pre-/periviable pregnancy loss or premature delivery.² Recognized risk factors for cervical incompetence include either congenital (müllerian anomalies, deficiencies in colaggen, elastine or connective tissue disease, and in utero exposure to diethylstibestrol) or acquired conditions (surgical trauma to the cervix, obstetric lacerations). Certain features of past (prior preterm birth, prior second trimester loss, induced abortion) or current pregnancy (cervical funnelling, multifetal pregnancy) may also increase the risk.^{1,3,4} The relative importance of such

factors varies between women and between subsequent pregnancies.^{1,3} It is difficult to ascertain the incidence of true cervical insufficiency, although data suggest estimates of 1% of women.²

The cervix is a relatively homogeneous structure, constituted mainly of collagen with very low cellular content, structural and physiologically different from the uterus. It's function, in normal pregnancy, is to retain the foetus in uterus initially and later easily allow passage and delivery, regaining retentive capacity subsequently.^{2,3}

Cervical insufficiency is classically defined as an asymptomatic (or painless) dilation and effacement of the cervix resulting in mid-trimester pregnancy loss or early preterm birth.² Since clinical criteria and diagnosis are difficult to establish and vary according to literature, its diagnosis is frequently made with a retrospective history of poor obstetric outcome.

As almost one in four women who have had a previous early spontaneous preterm delivery will recur in a subsequent pregnancy, several strategies to improve outcome have been proposed.

The cervix, regardless of whether the primary problem is, remains a logical target to intervein, even though it is unclear if cerclage will be beneficial to all women under this circumstances.²

Several cerclage techniques have arisen and evolved since its introduction in the 1950's. Its indications have been growing, from history of recurrent mid-trimester pregnancy loss to progressive cervical alterations and objective cervix dilation and effacement. This resulted in three types of cerclage: history-indicated (also known as prophylactic or primary), ultrasound-indicated (therapeutic or secondary) and physical exam-indicated cerclage (emergency, rescue or tertiary).^{4,5} Even for history-indicated guidelines criteria for performance varies widely worldwide, from history of 3 or more previous second-trimester pregnancy losses or preterm deliveries contrasting to history of 1 or more secondtrimester pregnancy loss related to painless cervical dilation in the absence of labour or placental abruption.⁴ Uniformity is required.

The aim of this study is to evaluate the maternal-foetal outcomes following prophylactic cervical cerclage at a level III perinatal care unit in Portugal.

METHODS

This is a hospital based, single-centre, retrospective study which reviewed data of all cases of women undergoing transvaginal history-indicated cervical cerclage from January 1st, 2008 to December 31st, 2017 at Centro Hospitalar Universitário do Algarve - Unidade Faro. Inclusion criteria met the American College of Obstetrics and Gynaecology (ACOG) Guideline indications for history-indicated cerclage: History of one or more second-trimester pregnancy losses related to painless cervical dilation and in the absence of labour or abruption placentae and/or prior cerclage due to painless cervical dilation in the second trimester. Excluding criteria consisted of follow up loss. Once the women were identified, all hospital records were obtained from the time of admission for cerclage placement to delivery, as well as data from previous pregnancies when available. Data on demographics, personal history, obstetric history, pregnancy, delivery and neonatal period were obtained. The primary outcome was gestational age <37weeks at birth. Secondary outcomes included hospital readmission during pregnancy, maternal morbidity, neonatal intensive care unit (NICU) admission and neonatal overall morbidity.

Statistical analysis

Data were analysed with IBM SPSS Statistics 23. Descriptive frequencies were used to present the results.

RESULTS

A total of 12 interventions (corresponding to 9 women) met the criteria and were involved in the study. Mean maternal age at the time of first cerclage was 27.6 years ± 4.39 ; when including recurrent cerclages, mean gestational age was 28.25 years ± 4.90 . At the time of first cerclage women had a mean of 2.44 previous gestations, a mean parity of 1.11 and only 50% of women had a previous live child (a mean of 0.78 live children) 87.7% of these being born preterm. The presence of known risk factors for cervical insufficiency was found in a minority (previous dilation and curettage and conization); no medical co-morbidities were identified. Demographics, obstetric history and risk factors are summarized in Table 1.

Table 1: Demographics, obstetric history and risk factors.

Age at first cerclage (years) (n= 9)	
Mean±SD	27.56±4.39
Median	28
Range	20-34
Age (years) (n= 12)	
Mean±SD	28.25 ± 4.90
Median	28
Range	20-38
Gravidity, mean (range)	2.44 (1-4)
Parity, mean (range)	1.11 (1-4)
Reproductive history	
Prior preterm birth (%)	83.4
Prior spontaneous abortion (%)	65.6
Risk factors for cervical insufficiency	
Cervical dilation for curetage (%)	22.2
Conization (%)	11.1

n: number of cases; SD: standard deviation; %: percentage.

In all cases McDonald technique was performed. Mean gestational age at hospital referral was 12.1weeks±2.76 with mean cervical length 35.1mm±5.13. At the time of cerclage placement, mean gestational age was 16.1weeks±0.51 and mean cervical length 27.5mm±8.70. Hospital admission lasted on average 10.7 days. Four hospital readmissions occurred: 3 for threatened preterm labour and 1 for falsely assumed beginning of labour.

Table 2: Cerclage placement and removal and foetal and maternal outcomes.

Gestational age at cerclage placement (weeks)	
Mean±SD	16.1±0.51
Range	15-17
Cervical length at placement (mm)	
Mean±SD	27.5 ± 8.70
Range	10-38
Gestational age at cerclage removal (weeks)	
Mean±SD	36.9±0.30
Range	36-37
Time from removal to delivery (days)	
Mean±SD	14.5±9.35
Range	1-28
Live birth (%)	100
Weight at birth (g) Mean±SD	3440.8±496.9
Apgar index >7 at min 5 (%)	100
Neonatl jaundice (%)	33.3
Need for phototherapy (%)	11.1
NICU admission	0
Maternal complications	0

SD: standard deviation; %: percentage; NICU = neonatal intensive care unit.

Removal of cerclage was made in ambulatory environment in 83.3% of cases. Mean gestational age at cerclage removal was 36.9 weeks and at delivery was 38,9 weeks, with average time between cerclage removal and labour being 14.5 days (range 1-28 days). Spontaneous onset of labour occurred in 75% of women, the remaining cases corresponding to induction of labour due to gestational diabetes or prolonged pregnancy. Delivery occurred vaginally in 83.4%. Caesarean section was performed in one case second to premature detachment of normally inserted placenta (PDNIP). Data on cerclage placement and removal and foetal and maternal outcomes are summarized in Table 2. No cases of preterm birth or stillbirth occurred. Neonates were female in 63.6% of case, weighted 3440.8g in average and Apgar index was above 7 at 5 minutes in all cases. Neonatal jaundice was observed in 25% of newborns, but no cases of respiratory distress, sepsis or major malformations were described. No cases were reported of foetal admission to NICU. There were no maternal complications.

After the intervention pregnancy, 100% of women had a live child, resulting in a mean number of live children of 1.58.

DISCUSSION

This analysis of hospital data reports a high rate of success after history-indicated cervical cerclage.

Similar results were found in the literature regarding history-indicated elective cerclage, achieving prevention of second trimester loss (STL) or preterm birth (PTB) in two thirds of low-risk groups (defined by one or two prior STL or PTB or by one previous successful cerclage).⁶⁻⁸

In this study one limitation may result from indication criteria. The indication for cerclage was at least one previous second trimester spontaneous pregnancy loss where no other cause was identified, conflicting with some recommendations were history of more pregnancy losses is required, hampering comparison.⁴ This fact may falsely improve outcomes, including women with less severe incompetence or better a priori outcome. However, cerclage was offered with these criteria to avoid a repeated miscarriage, weighting the psychological damage of a subsequent pregnancy loss.

In alternative, more strict criteria may result in very few eligible patients for history-indicated cerclage, thus decreasing the number of women benefiting from the procedure.⁹

Another limitation of this study is the reduced number of cases, either due to unregistered or lost cases from older registries or to low rate of cerclage proposal due to medical lack of confidence on ensuring the procedure. This limitation makes it inappropriate to generalize and over interpret the findings.

Since prematurity is a major cause of neonatal morbidity and mortality, with significant costs, a large number of interventions have been proposed for its prevention, some of which non-invasive. More recent systematic reviews and meta-analysis have compared progesterone, cerclage and pessary in decreasing preterm birth, with controversial results: one concluding that vaginal progesterone was the only intervention with consistent effectiveness for preventing preterm birth in singleton atrisk pregnancies whereas another defends that cervical cerclage showed clear benefit for women with singleton pregnancy and high risk of PTB.^{10,11} These alternatives were not considered for our patients as progesterone use in this context is a very recent recommendation and pessary is not available in our country.

CONCLUSION

Debate persists regarding strategies for decreasing preterm birth. Our results may show an over inclusion of women, when using only one prior pregnancy loss, resulting in unnecessary procedures. But, on the other hand, no maternal complications occurred. Thus, cervical cerclage might be an effective method to lower prematurity if provided for appropriately elected women, emphasizing the importance of re-evaluating the criteria for prophylactic cerclage.

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