Prediction of term and preterm parturition and treatment monitoring by measurement of cervical cross-linked collagen using light-induced fluorescence

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One of the keys to treating preterm labor is the early detection of changes indicating the onset of parturition.

Recently, we have developed a non-invasive method for the objective evaluation of the status of the cervix, where changes in collagen content of the cervix can be detected using an optical system and light-induced autofluorescence (LIF). This system measures the collagen fluorescence in the cervix as an indirect estimate of collagen concentration. Studies of pregnant women during the past few years support the use of this technique.

Cervical function and assessment

The cervix is composed of smooth muscle (ca. 10%) and a large component of connective tissue (90%) consisting of collagen, elastin, and macromolecular components, which make up the extracellular matrix (1–3). Many biochemical and functional changes occur in cervical connective tissue at the end of pregnancy (4–7). This process, termed cervical ripening, results in softening, dilatation, and effacement of the cervix. Ripening is required for appropriate progress of labor and delivery of the fetus.

Fluorescence spectroscopy of collagen

Fluorescence spectroscopy is a widely utilized research tool in the biosciences, primarily because of the amount of information that it can reveal in terms of molecular and physical states (8–14). We have used this methodology recently to evaluate the cervix during gestation.

The collascope and measurement of cervical ripening

Collagen gives characteristic fluorescence whose maximum is around 390 nm. The intrinsic fluorophor is believed to be pyridinoline, which is considered one of the major crosslinks within the primary structure of collagen fibrils (10, 15, 16). In our initial investigations, measurements were obtained from the serosal surface of the medium band of the cervix of rats in vivo. The results showed a decrease in fluorescence intensity decrease in the later gestational days and at parturition corresponding to the decrease in collagen. We also found a drop in collagen fluorescence intensity in rats treated with the anti-progesterone compound RU 38.486 and which delivered prematurely.
Further studies of rats

With the collascope, we were able to measure the fluorescence signal from the cervix in anaesthetized rats (17). The advantage to this technique is that one can follow cervical changes longitudinally in the same animal under a variety of conditions and treatments (Fig. 1).

In the postpartum period, the fluorescence gradually increased from the low value observed during delivery (Fig. 1). These results demonstrate the progressive decline in fluorescence during pregnancy to reach low values during delivery; these findings correlate well with cervical resistance as measured by the slope of stress–strain curves and a decline in cervical collagen content in electron micrographs of ripened versus unripe cervix. In addition, we examined rats at various times prior to and during preterm labor induced with the anti-progesterone onapristone. This study showed that ripening occurred with anti-progesterone treatment and that R5020, a progestin agonist, prevented ripening and preterm birth.

We conclude from these studies that the collascope can be used as a non-invasive tool to measure changes in cervical collagen content of the cervix under a variety of conditions. Results of these measurements correspond with known physiological changes in the cervix during pregnancy.

Studies of humans

We have also initiated human studies with the Collascope (17). Non-pregnant, pregnant, and postpartum human volunteers were recruited for the study. The first step was to establish a longitudinal distribution profile according to the weeks of gestation and postpartum. The cervical external os was gently wiped with rayon-tipped proctoscopic swabs prior to measurements being made. The measuring site was selected at the 12 o’clock position. Fluorescence decreased...
progressively during the final 15 weeks of pregnancy (Fig. 2). So far, several hundred patients (including nonpregnant patients) have been recruited. Several of the subjects have been measured two or three times during their pregnancy and postpartum. The results show a gradual decrease of the fluorescence as pregnancy approaches term followed by a slow recovery during the postpartum period (17–19). Recent studies demonstrated that the collascope could be used for the monitoring of the effect of induction agents on cervical ripening (20).

In view of the important role of collagen fibers and their turnover in the process of cervical function during pregnancy, the light-induced autofluorescence of cervical collagen could be a useful tool for evaluating cervical status and monitoring treatment strategies.

References


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