CAMPYLOBACTER FETUS SUBSP. FETUS ENUMERATION ON DIFFERENT MEDIA

CAMPYLOBACTER FETUS SUBSP.FETUS SKAITA NOTEIKŠANA DAŽĀDĀS BAROTNĒS

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ANOTĀCIJA. Darba mērķis bija noteikt *Campylobacter fetus* subsp. *fetus* augšanu atšķirīgās barotnēs dažādās atmosfērās. Pētījumā izmeklējamā kultūra tika iegūta no references celma ATCC 27374, kas inkubētas - 48 stundas 37±0.5°C mikroaerobā vidē. Rezultāti rāda, ka *Campylobacter fetus* subsp. *fetus* vislabāk aug uz Kolumbijas asins agara. Izvēloties selektīvas barotnes, kas paredzētas tieši *Campylobacter fetus* subsp. *fetus* izolēšanai, mCCDA un NA, labāku baktēriju augšanu novēro uz mCCDA barotnes. Datu statistiskā analīze tika veikta pēc t-testa ar ticamību 95%.

KEY WORDS: Campylobacter, fetus, recovery, media, enumeration

INTRODUCTION. Campylobacter fetus is gram-negative, curved or spiral, microaerophilic bacteria that cause a disease in cattle characterized primarily by sepsis, late embryonic death, infertility, a protracted calving season, and occasionally abortion. The aim of the study was to determinate Campylobacter fetus subsp. fetus recovery rate on different isolation media in different atmospheric conditions.

MATERIALS AND METHODS. Reference strain of Campylobacter fetus subsp. fetus ATCC 27374 was used. After primary incubation on two (sample A and B) Columbia blood agar (CA) (Oxoid; Basingstoke, Hampshire, UK) plates 10 µl of Campylobacter fetus subsp. fetus was transfered into 10ml of petone water (Oxoid) and vortexed until sample was disolved comletly to acquaire dilution 1:1000 or 10⁻³. Following that serial dilution 10⁻⁴ till 10⁻⁸ was prepared. After serial dilution 1ml of 10⁻⁷ and 10⁻⁸ dilution was transferred to two plates of Columbia blood agar (Oxoid), modified charcoal cefoperazone deoxycholate agar (mCCDA) (Oxoid) with Campylobacter selective supplement (Oxoid), nutrient agar I (NA) (Sifin; Berlin, Germany) with added Campylobacter growth and selective supplement (Sifin), and potato starch (MilliporeSigma; USA) 1.5 g/11. All plates were placed in anaerobic jars (Oxoid) and in order to provide microaerobic growth conditions CampyGen 3,51 (Oxoid) gas pack was added to the jars. Additionally, plates with Campylobacter fetus subsp. fetus with dillution 10⁻⁷ on mCCDA and CA was placed in candle jar with lit candle to reduce O₂ level. All plates were incubated for 48 h at 37±0.5 °C. After incubation enumeration of bacterial colonies was performed. Statistical analysis (t-test) was performed to determine statistically significant differences at 95% confidence level.

RESULTS. After 48 h of incubation in 37±0.5 °C on CA plates in sample A and B 14.87*10⁷ (CFU/ml), 11.82*10⁸ (CFU/ml), and 15.80*10⁷ (CFU/ml), 11.93*10⁸ (CFU/ml) cells were recovered, respectively. On mCCDA in sample A and B 14.11*10⁷ (CFU/ml), 11.04*10⁸ (CFU/ml), and 14.32*10⁷ (CFU/ml), 10.58*10⁸ (CFU/ml) cells were recovered, respectively. On nutrient agar I (NA) in sample A and B 10.62*10⁷ (CFU/ml), 2.4*10⁸ (CFU/ml), and 11.21*10⁷ (CFU/ml), 6.76*10⁸ (CFU/ml) cells were recovered, respectively. In candle jar on CA 13.59*10⁷ (CFU/ml) and on mCCDA 0.63 *10⁷ (CFU/ml) were recovered. Significantly

higher (p<0.05) recovery rate was observed in CA compered to mCCDA and NA, as well in mCCDA compered to NA (p<0.05).

CONCLUSIONS. For *Campylobacter fetus* subsp. *fetus* cultivation CA provides highest recovery rate. In case of selective media mCCDA is more superior than NA.

Candle jar can be used to provide microaerophilic conditions for *Campylobacter fetus* subsp. *fetus* cultivation on CA.