

Title: TEFLON MEMBRANES DO NOT ELIMINATE HALOTHANE INTERFERENCE WITH TRANSCUTANEOUS OXYGEN ELECTRODES

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**Introduction:** Continuing experience and technical improvements in transcutaneous oxygen (TC-O<sub>2</sub>) electrode systems have encouraged continuous tissue oxygen monitoring during general anesthesia. Since all commercially available TC-O<sub>2</sub> systems are polarographic and use a Clark electrode, however, they are subject to interference by gaseous molecules which can be reduced at the cathode. Careful choice of electrode membrane material and thickness could minimize interference by anesthetic gases. This study tested the concept that substituting Teflon for polypropylene membranes would eliminate halothane interference, previously described<sup>1</sup>.

**Methods:** Two Narco-AirShields (Hatboro, PA) Model TC-075-1 Transcutaneous Monitor Systems were used for this *in vitro* study. Duplicate electrodes each with a 25  $\mu$ m diameter platinum cathode and a 5.5 mm silver anode were used, one probe covered with a 20  $\mu$ m polypropylene membrane and the other with a 25  $\mu$ m Teflon PTFE membrane, both supplied by the manufacturer. Membranes were separated from the glass-covered electrode by a thin film of electrolyte solution. Electrode polarization was maintained at 600 mV and temperature at 44.4-44.5°C, with electrodes mounted in a test block and exposed to the same gas source simultaneously. Prior to each test, electrodes were polarized, zeroed, and calibrated with air after showing stability for a minimum of 24 hours. Gas mixtures were provided at flow rates of 1-3 l/min by an anesthesia machine with oxygen analyzer mounted at the fresh gas outlet. Indicated TC-O<sub>2</sub> readings were taken directly from the digital display of each monitor.

**Results:** Exposure of the polypropylene membrane system to halothane produced a rapid rise in indicated TC-O<sub>2</sub> despite constant oxygen partial pressure within 20, 10, or 5 minutes after beginning 0.5%, 1%, and 3% halothane, respectively (figure). By 3 hrs., a rise of 25 mmHg was seen for 0.5%, 90 mmHg for 1%, and 300 mmHg for 3% halothane, all in 21% O<sub>2</sub>, balance N<sub>2</sub>O.

No change in indicated TC-O<sub>2</sub> was seen with the Teflon system after 2 hours of 0.5% halothane in N<sub>2</sub>O-O<sub>2</sub>. However, TC-O<sub>2</sub> rose by 25 mmHg after 30 minutes of 3% halothane and by 6 mmHg after 120 minutes of 1% halothane in 21% O<sub>2</sub>, balance N<sub>2</sub>O.

Neither enflurane nor isoflurane at concentrations of 1 or 3% produced a significant discrepancy between the indicated TC-O<sub>2</sub> values when delivered in 21% O<sub>2</sub>, balance N<sub>2</sub>O, and TC-O<sub>2</sub>, varied less than 10 mmHg from the calculated P<sub>O<sub>2</sub></sub> of the test gas mixture at all times.

**Discussion:** Clinically useful transcutaneous oxygen monitoring during general anesthesia requires that indicated TC-O<sub>2</sub> values reflect tissue P<sub>O<sub>2</sub></sub>. Although careful attention to polarization voltages appears to have eliminated TC-O<sub>2</sub> artifact caused by N<sub>2</sub>O, our data indicate that halothane interference can give falsely high TC-O<sub>2</sub> readings even with Teflon PTFE membranes. These *in vitro* findings suggest that properly designed TC-O<sub>2</sub> systems may be used with confidence during general anesthesia with nitrous oxide, enflurane, or isoflurane. Sustained exposure to halothane in clinically-used concentrations at commonly encountered P<sub>O<sub>2</sub></sub> values, however, may produce falsely elevated indicated TC-O<sub>2</sub> when either Teflon or polypropylene membranes are used.

#### Reference:

1. Rafferty et al.: *In vitro* evaluation of a transcutaneous CO<sub>2</sub> and O<sub>2</sub> monitor: The effects of nitrous oxide, enflurane, and halothane. *Med. Instr.* 15: 316, 1981.

#### Figure:

