A204

PROPOFOL EFFECTS ON EEG AND RELATION-Title: SHIP WITH PLASMA CONCENTRATION DURING

NEUROSURGERY

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The goal of this study was to assess the relationship between propofol plasma concentration (PPC), EEG activity and cardiovascular (CV) effects1 during neurosurgery.

The protocol was approved by the ethics committee and informed written consent was obtained from 7 nonpremedicated patients (48.5±6.6 year, weighing 64.9±1.9 kg) with normal preoperative EEG, undergoing brain biopsies. EEG was recorded continuously from 4 electrodes (2 frontals, 2 occipitals). Anesthesia was induced with propofol 2 mg/kg i.v.; dextromoramide 0.05 mk/kg i.v.; vecuronium 0.01 mg/kg i.v. After intubation, anesthesia was maintained with propofol 6 mg/kg/h. The rate of infusion was increased by increments of 2 mg/kg/h until burst suppression was obtained (EEG stage I, maximum dose 14 mg/kg/h). According to the EEG changes, blood samples for PPC were collected during stages I to III. In addition, ECG (lead II), blood pressure and ETCO2 were monitored prior to and

during surgery.

Duration of anesthesia was 113±11.5 min, propofol dose was 8.93±1.03 mg/kg/h. Results are presented in Tables 1 and 2. Burst suppression required PPC of 5.02 mcg/ml.

Our data indicate that there is a positive relationship between PPC and EEG activity and that CV changes are minimal during propofol anesthesia. These results suggest that propofol may be a useful anesthetic during neurosurgery.

References

1. Anesth Analg 68:177-181, 1989.

Table 1. EEG Description of the Different Stages

Stage Description
I Burst suppression: 10-12 Hz or 1-4 Hz, low amplitude and electrical silence during 4-6 sec. Periodic activity: 10-12 Hz, low amplitude and 1-4 Hz low amplitude. 1-4 Hz moderate amplitude achieved with 12-15 Hz low amplitude.

Table 2. Data Function of EEG Stage

EEG	PPC	HR	MAP	RATE
	μg/ml	bts/min	mmHg	mg/kg/min
	Ö	82±6	113±6	Ō
I	5.09±0.64	69±3	92±3*	10.7±1.1
II	3.86±0.77	68±4	93±5*	2.7±1.2
III	1.60±3.0	76±4	109±5	0 .
	II	μg/ml 0 I 5.09±0.64 II 3.86±0.77	μg/ml bts/min 0 82±6 I 5.09±0.64 69±3 II 3.86±0.77 68±4	μg/ml bts/min mmHg 0 82±6 113±6 I 5.09±0.64 69±3 92±3* II 3.86±0.77 68±4 93±5*

Mean ± SEM; ANOVA one way; * P < 0.05 vs control

TITLE:

CEREBRAL BLOOD FLOW REACTIVITY

TO CO2 DURING HALOTHANE OR ISOFLURANE ANESTHESIA FOR CAROTID ENDARTERECTOMY

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Decreasing PaCO₂ during temporary vascular occlusion may have a beneficial influence on redistribution of CBF to ischemic brain.1 This study compared the effects of isoflurane (ISO) and halothane (HAL) on cerebral blood flow (CBF) reactivity to changes in PaCO₂ during carotid endarterectomy.

METHODS. After obtaining institutional approval and

informed consent, 14 patients, mean (\pm SE) age 68 \pm 2 received either ISO (n=12), 0.75% or O₂/N₂O, 1:1 or HAL (n=7), 0.5% in O2 / N2O, 1:1. Measurements were made during the period of temporary bypass shunting. Global CBF was measured using i.v. 133-Xe.² After a baseline CBF measurement, the PaCO₂ was elevated 8-10 mmHg by addition of CO2 to the inspired gas mixture and CBF was again measured. Data are expressed as mean \pm SEM.

RESULTS. By ANOVA, there was a significant effect of PaCO₂ to increase CBF but no difference between anesthetics, as shown in Fig. 1. There were no significant differences in mean blood pressure, temperature or heart rate between groups during the 2 measurements. No cerebral ischemia was detected by EEG. There was no difference (P = 0.913) in mean CBF reactivity to

changes in PaCO2 (ml/100g/min/mmHg) between ISO (1.7 ± 0.5) and HAL (1.8 ± 0.4). These values correspond to a relative 4 and 5 %-change in CBF per mmHg, respectively. The extrapolated CBF (ml/100g/min) at PaCO₂ = 40 mmHg tended (P = 0.126) to be higher for HAL (41 ± 4) than ISO (32 ± 3).

DISCUSSION. ISO increases the slope of feline CBF responsiveness to changes in PaCO2 in the hypocapnic range relative to HAL.³ In elderly patients undergoing carotid endarterectomy with relatively low maintenance concentrations of HAL or ISO in N2O there was no significant difference in their effects on CO₂ reactivity in mildly hypo- to normocapnic ranges. REFERENCES

- Anesthesiology 70:288-292, 1989
- Anesthesiology 71:863-869, 1989
 Anesthesiology 65:462-467, 1986

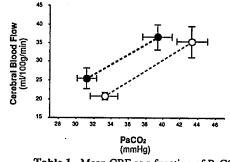


Table 1. Mean CBF as a function of PaCO2 for ISO (open circles) and HAL (closed circles).