Vasomotor Effects of Dimethyl Sulfoxide on Cat Cerebral Arteries In Vitro and In Vivo

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SUMMARY We studied the direct vascular effects of dimethyl sulfoxide (DMSO) in isolated middle cerebral arteries and on pial arteriolar caliber after subarachnoid perivascular microinjection in chloralose-anesthetized cats, and on brain retraction in cats given DMSO intravenously. DMSO did not constrict isolated cerebral arteries at any of the concentrations studied (10^{-10} to 4 \times 10^{-1} M). In middle cerebral arteries precontracted with potassium, 5-hydroxytryptamine, prostaglandin F_2 alpha, or with mechanically raised tone, DMSO at concentrations of 10^{-10} to 10^{-2} M had no significant effects; at concentrations greater than 10^{-2} M, DMSO consistently relaxed the arteries, probably because of the hyperosmolarity of the bathing solution. Microapplication of DMSO (10^{-10} to 10^{-2} M) around pial arterioles on the cortical surface did not change arteriolar caliber significantly. Higher concentrations of DMSO (1%) increased arteriolar caliber by 56 \pm 4\% (p < 0.001), probably as a consequence of solution hypertonicity. DMSO did not modify in vivo cerebrovascular responses to alterations in perivascular potassium ion concentrations. Intravenous administration of DMSO did cause obvious brain shrinkage. These data provide no support for the view that direct cerebral vascular effects play a major role in the clinical efficacy of DMSO, but are consistent with the hypothesis that DMSO's ability to lower intracranial pressure derives from its osmotic effect on cerebral tissue.

DIMETHYL SULFOXIDE (DMSO) has been reported to reduce morbidity and mortality in some but not other studies of cerebral infarction, pressure-induced ischemia, and trauma, and to lower traumatic intracranial hypertension and improve cerebral perfusion in humans and animals. In our current clinical trial of the effect of DMSO on raised intracranial pressure (ICP) after severe head injury, ICP was lowered within minutes after intravenous (IV) administration of DMSO in some patients (unpublished data). The mechanism of action of DMSO is unknown, although hypertonicity and direct cerebrovascular effects are two possible hypotheses. In this study, we investigated both the direct vasomotor effects of DMSO on isolated feline middle cerebral arteries in vitro and alterations resulting from microapplication of DMSO around individual pial arterioles on the cortical surface of anesthetized cats.

Materials and Methods
Response to Middle Cerebral Pial Arteries in Vitro

These experiments were conducted using techniques described previously. Six cats were anesthetized with 5% halothane, exsanguinated, and decapitated. The middle cerebral arteries were dissected and placed in cold Krebs'-Ringer's buffer solution (composition in mM: NaCl, 118; KCl, 4.5; MgSO_4 \cdot 2H_2O, 1.0; KH_2PO_4, 1.0; NaHCO_3, 25.0; CaCl_2 \cdot 2H_2O, 2.5; glucose, 6.0). Sections of artery 5 to 6 mm long were mounted on two L-shaped metal holders, one of which was attached to a force-displacement transducer to allow recording of the circular tension of the vessel to be measured, and placed in 50-ml organ bath to which various concentrations of DMSO could be added. The buffer solution was maintained at 37°C by an outer warming jacket and at pH 7.39 ± 0.01 (mean ± SEM) by continuous aeration with 5% CO_2/95% O_2. Initially, a 400-dyne load was placed on the vessel; during a 2-hour accommodation period, adjustments were made to maintain this tension, and the buffer solution was changed every 15 minutes. After a 15-minute period during which no further adjustments were needed to maintain the 400-dyne load, each vessel was tested with prostaglandin F_2 alpha (PGF_2 alpha, 2.5 \mu M). Vessels that did not constrict were excluded from further study. To accentuate the vasodilatory properties of DMSO, vessel tone was increased in some experiments by adding PGF_2 alpha (2.5 \mu M) or 5-hydroxytryptamine (5-HT, 0.3 \mu M) to the buffer, or by replacing the NaCl in the Krebs'-Ringer's solution with KCl to obtain a potassium concentration of 127 mM.

Log concentration-response curves were obtained by cumulative addition of DMSO to achieve bath concentrations of 10^{-10} to 4 \times 10^{-1} M (about 7.5 \times 10^{-10} to 3% DMSO by volume). DMSO concentrations that produced the half maximal response (EC_{50}) were calculated, as were the maximal responses (E_{max}) in dyne), for vessels with mechanically or pharmacologically increased tone.

Vasomotor Response of Pial Arterioles in Vivo

These investigations were performed using a previously described technique. Thirteen cats of either sex weighing 2 to 5 kg were anesthetized with an IV
infusion of alphaxalone, 9 mg/kg, and alphadolone, 3 mg/kg (Saffan, Glaxovet Ltd., Middlesex, England), intubated, and ventilated by intermittent positive pressure with a 33% N2O/67% O2 mixture. Anesthesia was maintained by periodic IV infusion of a 1% alphachloralose solution. Exhaled gases were monitored continuously with an infrared CO2 analyzer; arterial PO2, PCO2, and pH were measured periodically so they could be maintained within a physiologic range (PO2 = 172 ± 28 torr; PCO2 = 30 ± 2 torr; pH = 7.34 ± 0.04). The head of the cat was placed in a stereotaxic frame, and a 2 × 3-cm left parietal craniotomy was made with a dental drill under continuous saline irrigation. The dura was covered with a 1.5 cm layer of circulating mineral oil (38°C) and then opened under magnification (Bausch & Lomb stereomicroscope with zoom lens) and cold lighting from fiberoptic sources (Schott). Arteriolar calibers were measured by a standard image-splitting technique.13

Shortly before use, DMSO (Rimso, Research Industries, Salt Lake City, Utah) was dissolved to various concentrations (0.0001% to 1%) in normokalemic mock cerebrospinal fluid (CSF) (composition in mM: Na+, 156; K+, 3; Ca++, 2.5; HCO3-, 12; Cl-, 152) or mock CSF with 10-mM K+ (Na+, 149; K+, 10; Ca++, HCO3-, Cl- unchanged) or 40-mM K+ (Na+, 129; K+, 40; Ca++, HCO3-, Cl- unchanged). One percent DMSO was prepared by diluting 1 ml of pure DMSO (1.1 g) to 100 ml by addition of mock CSF. All solutions were adjusted to pH 7.2 by aeration with 5% CO2/95% O2 and aspirated under a thin film of mineral oil into glass micropipettes with tip diameters of 8 to 10 μm.

One series of injections was made with mock CSF containing increased NaCl (composition in mM: Na+, 224; K+, 3; Ca++, 2.5, HCO3-, 12; Cl-, 220) to produce a total solution osmolarity similar to that of 1% DMSO in mock CSF. A micromanipulator was used to place the micropipettes into the subarachnoid space near a pial vessel. A minute amount (about 5 μl) of solution was injected over 10 to 15 seconds; changes in vessel diameter were measured during the next 3 to 6 minutes, by which time the vessels had returned to control diameters. Alterations were expressed as the percent change from the injection diameter and compared by analysis of variance14 with those induced by injection of mock CSF.

Intravenous Infusion of DMSO

To evaluate the effect of DMSO on arterial pressure and brain size, DMSO (1 g/kg in a 10% solution) was infused intravenously in 2 cats. A blunt metal probe was positioned perpendicular to and gently on the brain and periodically adjusted to document changes in the position of the brain surface. At the end of the 1-hour infusion, a second dose of DMSO (1 g/kg in a 20% solution) was given over 3 to 5 minutes. Changes in blood pressure and brain surface location were noted.

Results

Middle Cerebral Arteries in Vitro

At concentrations of 10^-5 to 10^-2 M, DMSO had little effect on the tone generated by middle cerebral arteries (n = 6) in vitro, although it produced slight relaxation at higher concentrations (10^-2 to 4 × 10^-1 M) (fig. 1). In vessels constricted with PGF2alpha (220 ± 100 dyne; n = 11), DMSO resulted in vasodilatation equivalent to 85% of the induced tone, but only at the highest concentrations studied. The EC50 was 1.08 ± 0.35 × 10^-1 M and the EAm was −490 ± 70 dyne. 5-HT caused cerebral artery constriction amounting to 890 ± 150 dyne (n = 7). The subsequent cumulative addition of DMSO resulted in vasodilatation equal to 60% of the induced tone; again, this effect occurred only at the highest concentrations. The EC50 was 0.94 ± 0.70 × 10^-1 M and the EAm was not different from that in vessels constricted with PGF2alpha (≈ 510 ± 70 dyne). The buffer solution containing 127 mM potassium provoked maximal vasoconstriction of the cerebral vessels (950 ± 300 dyne; n = 13). Once again, DMSO relaxed middle cerebral arteries, but only in high concentrations (>10^-2 M); the amount of relaxation (13% of the induced tone) was somewhat less than in arteries preconstricted with PGF2 or 5-HT. However, the EC50 value was similar to that described above (1.56 ± 0.36 × 10^-1 M), and the EAm was −230 ± 50 dyne.

Pial Arterioles in Vivo

Perivascular microinjections of artificial CSF had little effect on pial arteriolar caliber (mean response, 1.5 ± 2.4%; n = 18). In artificial CSF, DMSO in concentrations of 0.0001% to 0.5% produced no significant effect on arteriolar caliber, although at high concentrations (e.g., 1%), arteriolar caliber was increased significantly (fig. 2). Neither the cerebrovascular dilatations provoked by microinjections of moderately elevated potassium concentrations (10 mM) nor the constrictions provoked by microinjections of markedly elevated potassium concentrations (40 mM)
were modified by the presence of DMSO (0.1%, or $10^{-2}$ M) in the injectate (fig. 3).

**Osmolar Effects**

The mean osmolarity of the Krebs' solution — 285 mosmol/l, determined by freezing point depression — was not altered by DMSO in concentrations of $10^{-10}$ to $10^{-2}$ M, but it was increased significantly by concentrations greater than $10^{-2}$ M (fig. 4). This change in osmolarity correlated extremely well with the concentration at which vasodilatation was first observed in the cerebral vessels.

The presence of DMSO in concentrations less than 0.01% had no effect on the osmolarity of the mock CSF, but higher concentrations produced concentration-dependent increases in osmolarity (e.g., artificial CSF: 299 ± 4 mosmol; DMSO 1%: 486 ± 5 mosmol). To determine the contribution of hypertonicity to the vascular effects of DMSO in vivo, hypertonic solutions of CSF (454 mosmol) produced by increasing the NaCl concentration were administered perivascularly.

**Arteriolar Responses to Potassium**

Arthuriolar caliber increased by 56 ± 4% ($p < 0.001; n = 6$) compared with the standard isotonic artificial CSF, and the dilatation produced by NaCl hyperosmolality was greater than that produced by 1% DMSO (19 ± 5%, $p < 0.001; n = 16$), which had similar osmolarity.

**Intravenous Infusion of DMSO**

After the last perivascular microinjection in 2 cats early in the series, intravenous infusion of 10% DMSO (1 g/kg) caused the brain to shrink visibly and recede from the edge of the dura. Later, in another 2 cats, we measured the changes in brain surface location and mean arterial pressure (MAP) during infusion of DMSO. In the first cat, the brain retracted 0.7 mm within 15 minutes after the start of the infusion (0.25 g/kg DMSO delivered) and remained retracted even though blood pressure rose from 75 to 100 mm Hg. Within 2 minutes after a second dose of DMSO, given more rapidly, MAP rose from 75 to 138 mm Hg, and was accompanied by a slight transient increase in brain size (maximal retraction, 0.4 mm). Ten minutes after the second injection, the brain retracted further, to a maximum of 1.9 mm, despite continuing hypertension (MAP = 110 mm Hg). In the second cat, the brain retracted 1.1 mm within 10 minutes after the start of the infusion (0.25 g/kg DMSO delivered) and remained retracted even though blood pressure rose from 75 to 100 mm Hg. Within 2 minutes after a second dose of DMSO, given more rapidly, MAP rose from 75 to 138 mm Hg, and was accompanied by a slight transient increase in brain size (maximal retraction, 0.4 mm). Ten minutes after the second injection, the brain retracted further, to a maximum of 1.9 mm, despite continuing hypertension (MAP = 110 mm Hg). In the second cat, the brain retracted 1.1 mm within 10 minutes after the start of the infusion (0.16 g/kg DMSO delivered), even though MAP increased from 112 to 127 mm Hg. At 1 hour, retraction was 2.1 mm despite hypertension to 160 mm Hg. The second DMSO injection further retracted the brain to a maximum of 2.9 mm when the MAP was 125 mm Hg. Postinjection hypertension to 190 mm Hg caused a slight increase in brain size, but even then the brain surface was 1.2 mm below its preinfusion position.
Blood volume. DMSO infusions have been shown to increase cerebral blood flow and osmotic equilibrium is established within 15 minutes. In experimental brain injury and in head injury patients at our institution (unpublished data) and elsewhere, DMSO has lowered ICP, sometimes within minutes. The mechanism by which DMSO lowers ICP is unknown, but the rapid onset of action raises the possibility of direct cerebral vasoconstriction and consequent reduction in intracranial volume and pressure.

In our experiments, however, DMSO did not cause cerebral vasoconstriction in vivo or in vitro at any concentration studied. In vitro, DMSO in concentrations at or above 0.2% produced significant and progressively greater vessel dilation; in vivo, vessel dilation occurred only with 1% DMSO solutions. These concentrations might be achieved transiently in the blood during IV DMSO infusion, but because of its ability to cross tissue membranes, DMSO distributes rapidly throughout the body, and clinical and experimental doses used probably yield DMSO tissue concentrations of 0.1% to 0.3% or less. Thus, the DMSO concentrations that produced arterial dilatation in our study probably exceed clinically achievable levels.

DMSO did exert significant relaxant effects on the cerebral vasculature both in vitro and in vivo, but only at concentrations sufficient to elevate the osmolarity of the organ bath or the injectate. Hyperosmolarity in blood and tissue causes vasodilatation in the peripheral circulation (muscle, gut, skin), and clinical and experimental doses used probably yield DMSO tissue concentrations of 0.1% to 0.3% or less. Thus, the DMSO concentrations that produced arterial dilatation in our study probably exceed clinically achievable levels.

In conclusion, DMSO exerted no direct vascular effect that would contribute to its ability to lower ICP, although cerebrovascular dilatation consistent with an osmotic effect occurred at high concentrations of the drug that would be unlikely to occur clinically. Nonetheless, systemic administration of DMSO in cats produced brain shrinkage, which is consistent with its reported ability to lower ICP. These observations emphasize the need for further investigation of the mechanisms of action of DMSO, but its lack of potentially deleterious actions on the cerebral vasculature and its ability to lower ICP justify continued use of DMSO in rigorously controlled studies of human head injury.

Acknowledgments

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References

11. Tsuruda J, James HE, Camp PE. Werner R: Acute dimethyl sulfox-
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