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Autoradiographic Method for Quantitative Evaluation of the Blood-Brain Barrier Effects of Contrast Media

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Opening of the blood-brain barrier after intravenous injection of different contrast media has been investigated by a quantitative autoradiographic technique using 14C-aminoisobutyric acid (AlB) as the blood-brain barrier radiotracer. In this study, experiments were carried out in adult rats. Animals were injected intravenously with 2 ml/kg of the tested contrast medium and immediately afterward with the blood-brain radiotracer AlB. The following contrast media have been tested: diatrizoate 38%, ioxithalamate 38%, ioxaglate 38%, the nonionic product metrizamide 40%, and a new nonionic product P-297, 400 mg I/ml. Control animals were injected intravenously with saline 0.9% (2 ml/kg) before the injection of the tracer. The degree of blood-brain barrier opening was quantitatively assessed by calculating the capillary rate constant for blood-to-brain transfer of AlB (k) from the brain activity and the arterial integral for a 6 min experiment. Preliminary data seem to indicate that the intravenous injection of 2 ml/kg of a constant medium may produce a tiny opening of the blood-brain barrier. But, if this is so, this blood-brain barrier opening is of a very low magnitude in the normal brain and there are no obvious differences between the tested contrast agents injected intravenously.

Different methods have been developed to determine the effects of contrast media on the blood-brain barrier (BBB). This problem has been classically investigated by vital dyes [1, 2]. Then, several investigators have used smaller tracer molecules, particularly the radionuclides, [32P]-orthophosphate [3] and [125I]-diacetyl [4], in an attempt to develop a more sensitive index for assessing BBB alterations. Another method has been electron microscopy [5, 6]. But none of these techniques has permitted us to quantify the effects of contrast media on BBB and to compare them with an objective index.

Therefore, to investigate this problem we selected a new quantitative radiographic method recently developed by Blasberg et al. [7, 8]. The BBB marker is 14C-aminoisobutyric acid (AlB), which is a small synthetic neutral amino acid, not metabolized; it has been found to be a very suitable tracer for quantifying BBB opening of moderate magnitude. This molecule does not cross normal brain capillaries rapidly and is avidly taken up and concentrated by brain cells. Because of these properties a simple transport model may be applied. According to this kinetic model, it is possible to calculate an unidirectional blood-to-brain transfer constant for AlB: k, from the following simple equation:

\[ k = \frac{C_v}{\int C_p \, dt} \]

in which \( C_v \) equals the concentration of the labelled material in the brain at the end of the experimental period, evaluated by quantitative autoradiography, and \( C_p \) is the tracer's concentration in the plasma.

In the present study, we have used this technique to evaluate the effects of contrast media on the BBB after intravenous injection, as with computed tomographic (CT) scanning. Since the widespread use of CT, large volumes of iodinated compounds have been injected intravenously. Physicians would like to select the contrast material with the lowest neurotoxicity, and, at the same time, the highest CT opacification to delineate an eventual lesion after intravenous injection.

Materials and Methods

Experiments were carried out in adult rats which were injected intravenously with a contrast medium at a dose of 2 ml/kg. The eventual opening of the BBB was evaluated by injecting the radiotracer AlB immediately afterward. Table 1 shows the contrast media we have tested. Control animals received 2 ml/kg of saline 0.9%. Three kinds of contrast media were investigated: (1) ionic monomeric compounds like diatrizoate or ioxithalame, which are the most hypertonic; (2) ionic dimeric compounds like ioxaglate, which is much less hypertonic; and (3) nonionic compounds like metrizamide and a new product from Guerbet Laboratory, P-297.

<table>
<thead>
<tr>
<th>Test Compounds used in this Study</th>
<th>Generic Name</th>
<th>Trade Name</th>
<th>mg I/ml</th>
</tr>
</thead>
<tbody>
<tr>
<td>Diatrizoate</td>
<td>Radioselectan</td>
<td>380</td>
<td></td>
</tr>
<tr>
<td>Ioxithalame</td>
<td>Telebrix</td>
<td>380</td>
<td></td>
</tr>
<tr>
<td>Ioxaglate</td>
<td>Hexabrix</td>
<td>380</td>
<td></td>
</tr>
<tr>
<td>Metrizamide</td>
<td>Amipaque</td>
<td>400</td>
<td></td>
</tr>
<tr>
<td>P-297</td>
<td></td>
<td>400</td>
<td></td>
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</tbody>
</table>

Note.—Control animals received NaCl 0.9%.
The experimental procedure included the following steps: (1) insertion of catheters in the femoral vein and artery; (2) a bolus intravenous injection of the contrast medium, plus the tracer AIB (100 µCi [0.37 Bq]), serial arterial blood sampling for 14C-counting in plasma, and, 6 min after start, sacrifice of the animal and rapid freezing of the brain; (3) serial sectioning of frozen brain (20 µm) in a cryomicrotome at −20°C; (4) quantitative autoradiography; and (5) calculation of the blood-to-brain transfer constant for AIB (k) in the following cerebral structures: olfactory cortex, parietal cortex, striatum, occipital cortex, and white matter of pons.

Quantitative autoradiography requires the exposure on the same film (Kodak SB-5) of brain sections of the same thickness and of calibrated 14C-standards with different known 14C-concentrations. The optical density of standards is measured on the film and correlated to 14C-concentration; from this calibration curve, the 14C-concentration of a cerebral structure may be calculated after measurement of its optical density.

Results and Discussion

 Autoradiographs of 14C-AIB distribution within rat brain in control and experimental animals are presented in figures 1 and 2. In control animals (figs. 1A and 2A), we may observe that 14C-activity in cerebral structures is remarkably low, except in the area of choroid plexus. Compared with the control, 14C-AIB distribution seems to be slightly increased after the intravenous injection of Diatrizoate (fig. 1B) or Ioxaglate (fig. 1C). Autoradiographs from animals injected with nonionic contrast media (figs. 2B and 2C) also demonstrate a slight increase of 14C-AIB distribution. However, in all cases, the variation of 14C-AIB distribution pattern does not seem to be of a great magnitude after the intravenous injection of contrast media; no major difference is obvious between the tested contrast media.

Tables 2 and 3 summarize the preliminary quantitative data we have obtained. Until now, our data are limited to a few animals: three controls and 10 experimental animals injected by contrast media (two animals for each contrast agent).

For each animal, the blood-to-brain transfer constant for AIB (k) was calculated in five cerebral structures, as indicated on table 2. In this table, there are quite similar values of k for the different gray matter structures, and a relative lower value for white matter, which is probably due to differences in capillary density rather than differences in capillary permeability. Because of the similarity of k values in different gray matter structures, table 3 shows only the blood-to-brain transfer constant for AIB measured in the parietal cortex for each animal.

There were too few animals to evaluate the results statistically.
However, table 3 reveals that in control animals the blood-to-brain transfer constant for AIB was 5–10 U, while in animals injected with different contrast media values of k, in most of the cases were above the control values (in most animals, k, was greater than 10 U). On the other hand, the results in table 3 suggest there may not be significant differences of k, among the tested contrast media.

In summary, our data indicate that the intravenous injection of 2 ml/kg of a contrast medium may produce a tiny opening of BBB, as tested by the AIB transfer measurement. But, if it is so, this opening is of a very low magnitude in a normal brain, and there are no obvious differences between the tested media when they are injected intravenously. These findings may be helpful to further investigate problems of neurotoxicity of contrast media and CT enhancement after their injection. Blood-brain barrier opening after intravenous injection of contrast media does not seem to be the most important problem. What the contrast medium is doing to the central nervous system tissue and how it moves and diffuses through the parenchyma after crossing the BBB are probably of more interest.

REFERENCES

4. Golman K. The blood-brain-barrier: effects of non-ionic contrast media with and without addition of Ca²⁺ and Mg²⁺. *Invest Radiol* 1979;14:305–308