THE EFFECT OF THE FORMATION OF AN ARTERIOVENOUS FISTULA UPON BLOOD VOLUME 1

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The creation of a large peripheral arteriovenous fistula results in a series of related physiologic adjustments, including changes in cardiac output, intravascular and intracardiac pressures, blood flow to peripheral organs and urinary excretion of water and electrolytes (1–3). The changes in blood volume which occur under these circumstances have been the subject of controversy, partly because observations made before and after surgical extirpation of a fistula are necessarily complicated by the effects of a major operation and the patient’s convalescence.

Previous efforts to resolve this question have utilized protein-binding dyes such as T-1824 or Brilliant Vital Red to measure blood volume. However, the accuracy of T-1824 as a measure of total blood volume is open to question (4) and its validity as an index to the intravascular volume of plasma in certain conditions has been challenged (5, 6). For these reasons it was decided to reinvestigate the early effect of an experimental arteriovenous fistula upon blood volume, and to use labelled red cells as well as T-1824 in an attempt to measure by independent methods changes in the volume of red cells and plasma.

METHODS

Seven trained mongrel dogs, weighing 10 to 18 kilograms, were fed a standard meal diet to which raw meat and ferrous sulfate were added. After control measurements of blood volume in the unanesthetized, fasting state, an arteriovenous fistula approximately one cm. long was created surgically between the abdominal aorta and the inferior vena cava, halfway between the renal vessels and the bifurcation. From two to six weeks after the operation, the blood volume was remeasured.

Five cc. of the dog’s own blood were incubated at room temperature for 30 minutes with Pni. After labeling, the red cells were washed three times with cold saline and reinjected intravenously from a calibrated syringe. Supernatant saline from the last wash contained less than 2 per cent of the total radioactivity incorporated in the red cells. The injection of cells was immediately followed by the intravenous injection of 10 mgm. of Evans Blue dye (T-1824). Heparinized samples of arterial blood were drawn at five-minute intervals for 30 minutes. Radioactivity of whole blood was measured by the method of Reeve and Veall (7). Optical density of plasma diluted with an equal volume of saline was recorded at 610 λ using a Coleman Jr. spectrophotometer. Concentrations of radioactivity in blood and of T-1824 in plasma were plotted on a semilogarithmic scale against time and the straight lines of best fit were extrapolated to the times of injection (T0). Hematocrit of arterial blood was determined by spinning 100 mm. columns of blood in Wintrobe tubes in duplicate for one hour at 3000 rpm. in a centrifuge head of 15 cm. radius from the center of the spindle to the bottom of the hematocrit tube. A correction of 0.96 was applied for plasma trapping. The following calculations were used:

\[
P^{mw} = \frac{\text{Radioactivity injected}}{\text{Radioactivity of 1 cc. arterial blood at T0}}
\]

1) Total Blood Volume (TBV \(P^{mw}\))

\[
2) \text{RBC Volume (RBC \(P^{mw}\))} = (\text{TBV \(P^{mw}\) \times Arterial Hematocrit \times 0.96})
\]

3) \(T-1824\) space \(=\) \(\frac{T-1824 \text{ injected}}{\text{T-1824 in 1 cc. arterial plasma at T0}}\)

4) Total Blood Volume (TBV \(T-1824\)) \(=\) \(\frac{T-1824 \text{ space}}{1 - (\text{Arterial Hematocrit \times 0.96})}\)

Combined

5) Total Blood Volume (TBV comb) \(= (\text{RBC \(P^{mw}\)} + T-1824 \text{ space})\)

6) Total Body Hematocrit (TB Hct) \(=\) \(\frac{\text{RBC \(P^{mw}\)}}{\text{TBV comb}}\)

RESULTS

(Table I)

I. General

Following the operation, cardiac filling pressure was chronically elevated and the cardiac output increased. Cardiac output could not be increased

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### TABLE 1

<table>
<thead>
<tr>
<th>Days before or after fistula</th>
<th>Wt.</th>
<th>RBC vol.</th>
<th>TBV</th>
<th>Plasma vol.</th>
<th>TBV</th>
<th>TBV comb</th>
<th>Arterial hct</th>
<th>TB hct</th>
<th>TB hct</th>
<th>Edema/Ascites</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 - 6</td>
<td>kg.</td>
<td>cc.</td>
<td>cc.</td>
<td>cc.</td>
<td>cc.</td>
<td>%</td>
<td>%</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>+33</td>
<td></td>
<td>329</td>
<td>851</td>
<td>582</td>
<td>948</td>
<td>911</td>
<td>38.6</td>
<td>36.1</td>
<td>.944</td>
<td>0/0</td>
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<tr>
<td>2 - 14</td>
<td></td>
<td>11.1</td>
<td>324</td>
<td>885</td>
<td>901</td>
<td>894</td>
<td>36.7</td>
<td>36.2</td>
<td>.980</td>
<td>++/0</td>
</tr>
<tr>
<td>+20</td>
<td></td>
<td>11.3</td>
<td>322</td>
<td>906</td>
<td>700</td>
<td>1,022</td>
<td>35.6</td>
<td>31.5</td>
<td>.885</td>
<td>0/+</td>
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<tr>
<td>3 - 12</td>
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<td>10.9</td>
<td>477</td>
<td>1,037</td>
<td>620</td>
<td>1,097</td>
<td>46.0</td>
<td>43.5</td>
<td>.943</td>
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<td>11.1</td>
<td>358</td>
<td>980</td>
<td>935</td>
<td>1,293</td>
<td>39.0</td>
<td>27.7</td>
<td>.710</td>
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<td>12.7</td>
<td>340</td>
<td>904</td>
<td>636</td>
<td>976</td>
<td>37.6</td>
<td>34.8</td>
<td>.925</td>
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<td>+14</td>
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<td>13.0</td>
<td>320</td>
<td>1,033</td>
<td>803</td>
<td>1,123</td>
<td>31.0</td>
<td>28.5</td>
<td>.917</td>
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<td>17.8</td>
<td>556</td>
<td>1,572</td>
<td>1,168</td>
<td>1,724</td>
<td>35.4</td>
<td>32.3</td>
<td>.917</td>
<td></td>
</tr>
<tr>
<td>+14</td>
<td></td>
<td>17.3</td>
<td>566</td>
<td>1,788</td>
<td>1,470</td>
<td>2,036</td>
<td>31.7</td>
<td>27.8</td>
<td>.877</td>
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<td>6 - 3</td>
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<td>12.3</td>
<td>415</td>
<td>1,000</td>
<td>705</td>
<td>1,120</td>
<td>41.5</td>
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<tr>
<td>+42</td>
<td></td>
<td>12.5</td>
<td>490</td>
<td>1,311</td>
<td>970</td>
<td>1,460</td>
<td>37.4</td>
<td>33.5</td>
<td>.892</td>
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<tr>
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<td>14.8</td>
<td>652</td>
<td>1,265</td>
<td>837</td>
<td>1,489</td>
<td>51.6</td>
<td>43.7</td>
<td>.847</td>
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<tr>
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<td></td>
<td>15.5</td>
<td>565</td>
<td>1,542</td>
<td>1,137</td>
<td>1,702</td>
<td>42.1</td>
<td>33.2</td>
<td>.786</td>
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<tr>
<td>Mean - 6</td>
<td></td>
<td>13.1</td>
<td>442</td>
<td>1,073</td>
<td>731</td>
<td>1,173</td>
<td>41.1</td>
<td>37.7</td>
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<tr>
<td>+26</td>
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<td>13.2</td>
<td>424</td>
<td>1,192</td>
<td>973</td>
<td>1,397</td>
<td>36.0</td>
<td>30.2</td>
<td>.833</td>
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<tr>
<td>Δ</td>
<td></td>
<td>-18</td>
<td>+119</td>
<td>+242</td>
<td>+267</td>
<td>+224</td>
<td>-5.1</td>
<td>-7.5</td>
<td>-0.84</td>
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</tr>
<tr>
<td>p</td>
<td></td>
<td>0.5</td>
<td>0.05</td>
<td>0.01</td>
<td>0.01</td>
<td>0.01</td>
<td>0.02-0.05</td>
<td></td>
<td></td>
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</tr>
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</table>

It should be noted that these fistulas were much larger in proportion to body size than those usually seen clinically.

II. Total red cell volume (Figure 1)

After an arteriovenous fistula there was no consistent change in total red cell volume measured with radioactive red cells. The average change was - 18 cc., with variations from - 119 cc. to + 75 cc.

III. Plasma volume (Figure 2)

In contrast to the relative constancy of red cell volume, the volume of distribution of T-1824 expanded markedly in every instance, and by an average of 242 cc., or 32 per cent. That this reflected, at least in part, an actual increase in plasma volume following creation of an arteriovenous fistula is suggested by an average fall in arterial hematocrit of five millimeters, in the presence of an unchanged total red cell volume. In this small series, no correlation was apparent between the presence or absence of ascites or pitting edema and the degree of expansion of the plasma volume.

There were no consistent differences between
the slopes of disappearance of T-1824 in normal dogs and in the same dogs after creation of a fistula. Blue dye was not detected in samples of ascitic fluid removed 30 minutes after intravenous injection of the dye.

IV. Total blood volume and total body hematocrit

In every dog, total blood volume was increased several weeks after an arteriovenous fistula, whether this was calculated from the dilution of P32-tagged cells, from the T-1824 space and the arterial hematocrit, or by adding the directly determined red cell volume and the T-1824 space. This was a result of the increase in the volume of plasma. As others have noted (4), values for blood volume calculated from the dilution of blue dye were invariably higher than those obtained by the red cell technique. Furthermore, the expansion of blood and plasma volume after a fistula was significantly greater, both absolutely and percentagewise, when measured with T-1824, than when calculated from the directly measured, unchanged red cell volume and the fall in arterial hematocrit. Another way to express this is as a decrease in the ratio of total body hematocrit to arterial hematocrit. As seen in Figure 3, this ratio diminished in six of seven instances.

DISCUSSION

The sudden direct transmission of arterial pressure to the distensible great veins which occurs when an arteriovenous fistula is opened must increase the volume of blood contained in the venous tree, with simultaneous contraction of other portions of the vascular system. A progressive increase in total blood volume under these circumstances would mitigate the necessity for such a localized vascular contraction. In 1924, Holman reported that the volume of distribution of protein-bound dyes was increased in the presence of an arteriovenous fistula (11). Although these findings were confirmed in certain subsequent studies (12–16), they were not supported by others (17, 18). Warren, Elkin, and Nickerson reported that less than half of forty-one patients with an arteriovenous fistula had a significant fall in blood volume estimated from the T-1824 space following surgical repair of the defect (19). On the other hand, Schreiner, Freinkel, Athens, and Stone found a post-operative decrease in T-1824 space of more than five per cent in sixteen of nineteen patients studied, although no change in hematocrit was noted (20). Lillehei, Bobb, and Visscher found a marked increase in both T-1824 space and thiocyanate space after formation of an arteriovenous fistula in dogs (21).

Although total blood volume increased in every animal in the present study after creation of a fistula, this was accomplished chiefly by an expansion of the volume of plasma, without consistent changes in total red cell volume. Plasma volume increased despite an elevation in venous (and presumably capillary) pressure which would favor increased transudation of fluid out of the vascular system. Renal retention of salt and water, stimulated, perhaps, by inadequate arterial filling (2, 3), presumably plays an important role.
in the expansion of extracellular fluid and plasma which occurs under these circumstances. Mobilization of extravascular depots of protein may also play a part in establishing and maintaining the expanded plasma volume (20, 22).

The T-1824 space in five animals with fistulas enlarged to a greater extent than would have been predicted from the concomitant fall in arterial hematocrit, resulting in a decrease in the ratio total body hematocrit: arterial hematocrit. A similar decrease in this ratio is characteristic of patients with elevated venous pressure and edema secondary to congestive heart failure (23). Although interpretation of this finding may be difficult in the dog with intact spleen (24), three possible explanations might be suggested. First, it is possible that in a dog with a large fistula, the splenic reservoir of blood rich in red cells may contract and become smaller. Secondly, the decreased ratio might reflect a redistribution of blood between large and small vessels. This would require a larger proportion of blood to be contained in small vessels with low hematocrit in the animal with a fistula, in the face of obviously increasing engorgement of the great central veins. Finally, it is well known that protein tagged with Evans Blue dye circulates through perivascular lymph channels and extravascular protein depots, as well as inside the vascular system (25). It is conceivable that with the widened pulse pressure, elevated venous pressure and heightened rapidity of the circulation consequent to an arteriovenous fistula, the initial passage of dye into lymph channels or other easily accessible extravascular compartments might be somewhat accelerated and its calculated initial volume of distribution therefore expanded. The present experiments are consistent with such a supposition though they provide no direct evidence for it.

SUMMARY

Blood volume was measured in seven dogs, using P32-tagged red cells and T-1824, before and up to six weeks after a large fistula was created between the aorta and inferior vena cava. The following were noted:

1. No change in total red cell volume.
2. An expanded plasma volume, reflected in a lowered hematocrit and an increase in the volume of distribution of T-1824.
3. A decrease in the ratio of total body hematocrit to the hematocrit of arterial blood.

REFERENCES