Ringer-lactate solution as a priming fluid for the disc oxygenator

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Gadboys, Slonim, and Litwak (1962) reported what is now known as the homologous blood syndrome, which stimulated work on a substitute for blood in open-heart surgery, especially for the priming of the heart–lung machine.

The homologous blood syndrome was noted in experimental animals and consisted of hypotension on commencing cardiopulmonary bypass, with early thrombocytopenia, leucopenia, sequestration of blood elements, and subsequent pulmonary and renal problems. Gadboys et al. concluded that the changes were consistent with antigen–antibody reactions other than those due to ABO and Rh incompatibilities.

An obvious substitute was a colloid, such as a low molecular weight dextran (rheomacrodex) (Long, Sanchez, Varco, and Lillehei, 1961; Raison, 1962). Others have described using electrolyte solutions as partial primes and also as total non-blood primes.

Neptune, Bougas, and Panico (1961) reported the use of physiological saline; Zuhdi, McCollough, Carey, and Greer (1961), Cooley, Beall, and Grondin (1962), and Zuhdi, Carey, Sheldon, and Greer (1964) used 5% dextrose in water, but as this was physiologically hypotonic, Linder, Sakai, and Paton (1964) attempted to improve the solution by using varying concentrations of saline rather than water. They found that 5% dextrose in 0.22% saline gave the best results.

Neville, Faber, and Peacock (1964a) showed that Ringer-lactate solution (Hartmann's solution) caused minimal disturbances of acid/base and electrolyte homeostasis and seemed to be the most physiologically acceptable priming fluid available.

Total non-blood priming has required heart–lung machines with reasonably small priming volumes, and this has discouraged the use of this technique with the disc type of oxygenator, though Neville et al. (1964a) used the Kay-Cross oxygenator. Before changing to the Rygg-Kvysgaard disposable oxygenator experience was gained with haemodilution using the Guy's-Ross heart–lung machine with disc oxygenators, and this paper aims to show how practical, satisfactory, and economical this technique has proved to be.

TECHNIQUE

Twenty-two consecutive patients who were undergoing procedures requiring cardiopulmonary bypass were studied. Their ages ranged from 7 to 49 years, and body weights from 21 to 75 kg. Each patient received suitable doses of omnopon and scopolamine as premedication. Anaesthesia was induced with thiopentone (2.5%) and was maintained with nitrous oxide and oxygen, and tubocurarine. Oxygen containing 0.5% halothane was fed into the oxygenator, and during total body perfusion the lungs were kept inflated with 100% oxygen.

The Guy's-Ross rotating disc oxygenator was used (Ross, 1960). This requires 2 litres of priming fluid for flows up to 2.5 litres/minute and 3 litres when an additional oxygenator was added, allowing flows up to 4 litres/minute. Acid-citrate-dextrose (A.C.D.) blood was used in preference to fresh blood both in the oxygenator, when necessary, and for direct intravenous infusion throughout the surgery. Theoretically, blood which is four to five days old is desirable in order to minimize the body reactions due to white cells and platelets.

Haemodilution was carried out with Ringer-lactate solution, with the addition of 20 mEq sodium bicarbonate per litre of priming fluid in those patients who had a total non-blood prime. The average pH of Ringer-lactate is 6.5, and, in the absence of any blood, it was found necessary to counteract this with the bicarbonate (see Discussion). The degree of haemodilution was based on the estimated blood volume and the patient's haematocrit. The most accurate figure for estimating the blood volume proved...
to be 65 ml./kg. body weight (Wasserman, Yoh, and Raschoff, 1951). The volume of Ringer-lactate used in the prime was calculated to bring the packed cell volume (P.C.V.) to a figure of between 20 and 24%. Initially, the dilution used led to figures above this value, but the simplicity and safety of the technique led to increasing degrees of haemodilution (Table I) until, with suitably sized patients, a complete pump prime with Ringer-lactate was possible.

The patients were fully heparinized with 3 mg./kg. body weight and, in order to prevent dilution of heparin during perfusion, heparin was also added to the pump at the time of priming (30 mg./litre).

The perfusion was started on a partial basis using a small flow, initially—not more than 25% of estimated full flow until adequate mixing had occurred or until cooling was sufficient, if this was necessary. Full flows were based on a flow of 2-4 litres/m.² body surface at normal body temperature.

An estimate of P.C.V., by the microhaematocrit method, was carried out after 10 minutes, and this figure was always the lowest that was recorded, suggesting that adequate mixing had occurred within this period. The patient was initially perfused with a positive fluid balance of about 200 ml, and this was increased during perfusion to about 1 litre after an hour.

Arterial blood pressure was often low, and figures of 20 mm. Hg were not unusual during total perfusion. As soon as perfusion ended, the patient's red cell mass was restored to normal by infusing the fluid remaining in the pump. When only a partial non-blood prime was used, the volume of fluid infused was 1,000 to 1,800 ml., depending on calculations based on the patient's estimated blood volume and P.C.V. and on the P.C.V. of the perfusate at the end of bypass. Where no blood was used in the prime, the full contents of the extracorporeal circuit were returned to the patient. This was as much as 3,000 ml. when the double oxygenator was used.

The patient's blood volume was raised by the infusion of volumes of 200 ml, and a continuous record of the central venous pressure was available. This showed a small rise following each 200 ml, which settled within two minutes. The next 200 ml was then given. In this way the full volume could be given within 20 minutes, and the venous pressure at this point was usually little above normal.

Heparin was reversed, at the end of bypass, with protamine in a dosage of 4 mg./kg. body weight, no account being taken of the heparin initially put in the pump. A heparin–protamine titration was carried out 20 minutes after the protamine had been given, and suitable adjustment was made if necessary.

In the post-operative period, apart from blood, only 5% dextrose in water was given by intravenous infusion. This was discontinued as soon as possible. Loss of blood was measured during and after operation. Any loss in the preperfusion period was replaced with A.C.D. blood before perfusion to obviate any further reduction in P.C.V. during the perfusion.

INVESTIGATIONS

Before operation the plasma urea and electrolytes, full blood count, bleeding and clotting times, prothrombin time, plasma fibrinogen, P.C.V., and plasma-free haemoglobin (P.F.Hb) were measured. A 24-hour specimen of urine was collected and examined.

At the start of the operation a urinary catheter was passed and the urinary output was measured hourly for 24 hours. During this time, the excretion of urea and electrolytes was measured at eight-hourly intervals. In the succeeding two days, 24-hour specimens were similarly examined.

The P.C.V., Hb, and P.F.Hb were measured at 20-minute intervals during perfusion, one and three hours after perfusion, and on the next two days. Plasma electrolytes were measured one hour after perfusion and on the next two days.

At the end of perfusion the arterial pH, Pco₂, and standard bicarbonate were measured by the Astrup technique (Andersen, Engel, Jørgensen, and Astrup, 1960). When necessary, sodium bicarbonate was then given, a level of 20 mEq/l. being regarded as a definite indication.

RESULTS

These are summarized in Table I. All 22 patients survived and left hospital, seven of them having had a total non-blood prime.

The P.C.V. fell to its lowest level within 10 minutes of the start of perfusion, and then rose from 1% to 6% during the remainder of the perfusion. This rise was not related to the length of perfusion. Within one hour of the end of perfusion, the P.C.V. had returned to within normal limits. Haemoglobin estimations provided a parallel series of results. Owing to technical difficulties, only five patients had estimations of P.F.Hb. Haemolysis reached a maximum towards the end of perfusion, and the highest figure recorded was 300 mg./100 ml. This soon fell and approached normal levels after two days.

Three patients developed petechiae post-operatively.

At the end of perfusion, only four patients showed a base deficit sufficient to require sodium bicarbonate.

Plasma sodium and chloride showed a normal post-operative fall and remained within normal limits. The plasma potassium, however, fell below 4 mEq/l, in seven patients within 24 hours, and in 11 patients within 48 hours. This is the reverse of that expected in the metabolic response to trauma (Le Quesne, 1959).

Within 20 minutes of the end of perfusion a volume of perfusion fluid varying from 825 ml. (case 12) to 3,000 ml. (case 10) was returned to
Ringer-lactate solution as a priming fluid for the disc oxygenator

**TABLE I**

RESULTS IN 22 PATIENTS USING RINGER-LACTATE SOLUTION IN THE PRIMING FLUID

<table>
<thead>
<tr>
<th>Case</th>
<th>Sex</th>
<th>Age (yrs)</th>
<th>Weight (kg.)</th>
<th>Surface Area (m²)</th>
<th>Operative Procedure</th>
<th>Perfusion (min.)</th>
<th>Haemodilution (ml./kg.)</th>
<th>Lowest P.C.V. (%)</th>
<th>Lowest Temp. (°C.)</th>
<th>Priming Fluid</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>M</td>
<td>12</td>
<td>54</td>
<td>1-5</td>
<td>Aortic valvotomy (subvalvar)</td>
<td>54</td>
<td>37</td>
<td>30</td>
<td>28</td>
<td>31</td>
</tr>
<tr>
<td>2</td>
<td>M</td>
<td>26</td>
<td>69</td>
<td>1-9</td>
<td>Repair A.S.D. (secundum)</td>
<td>39</td>
<td>29</td>
<td>33</td>
<td>33</td>
<td>33</td>
</tr>
<tr>
<td>3</td>
<td>F</td>
<td>49</td>
<td>63</td>
<td>1-7</td>
<td>Aortic valvotomy</td>
<td>108</td>
<td>36</td>
<td>25</td>
<td>24</td>
<td>24</td>
</tr>
<tr>
<td>4</td>
<td>F</td>
<td>36</td>
<td>51</td>
<td>1-5</td>
<td>Aortic valvotomy</td>
<td>78</td>
<td>39</td>
<td>25</td>
<td>25</td>
<td>27</td>
</tr>
<tr>
<td>5</td>
<td>F</td>
<td>10</td>
<td>35</td>
<td>1-2</td>
<td>Pulmonary valvotomy</td>
<td>75</td>
<td>43</td>
<td>25</td>
<td>29</td>
<td>32</td>
</tr>
<tr>
<td>6</td>
<td>M</td>
<td>17</td>
<td>60</td>
<td>1-7</td>
<td>Pulmonary valvotomy. Repair V.S.D.</td>
<td>115</td>
<td>41</td>
<td>28</td>
<td>26</td>
<td>28</td>
</tr>
<tr>
<td>7</td>
<td>M</td>
<td>14</td>
<td>31</td>
<td>1-1</td>
<td>Repair V.S.D.</td>
<td>104</td>
<td>48</td>
<td>25</td>
<td>18</td>
<td>21</td>
</tr>
<tr>
<td>8</td>
<td>M</td>
<td>8</td>
<td>25</td>
<td>1-0</td>
<td>Pulmonary valvotomy</td>
<td>74</td>
<td>60</td>
<td>28</td>
<td>31</td>
<td>32</td>
</tr>
<tr>
<td>9</td>
<td>F</td>
<td>10</td>
<td>38</td>
<td>1-3</td>
<td>Repair V.S.D.</td>
<td>85</td>
<td>40</td>
<td>26</td>
<td>22</td>
<td>28</td>
</tr>
<tr>
<td>10</td>
<td>F</td>
<td>14</td>
<td>48</td>
<td>1-4</td>
<td>Pulmonary valvotomy. Repair V.S.D.</td>
<td>75</td>
<td>62</td>
<td>22</td>
<td>21</td>
<td>27</td>
</tr>
<tr>
<td>11</td>
<td>F</td>
<td>16</td>
<td>42</td>
<td>1-3</td>
<td>Repair V.S.D.</td>
<td>60</td>
<td>47</td>
<td>22</td>
<td>28</td>
<td>30</td>
</tr>
<tr>
<td>12</td>
<td>M</td>
<td>7</td>
<td>21</td>
<td>0-9</td>
<td>Aortic valvotomy</td>
<td>55</td>
<td>60</td>
<td>26</td>
<td>33</td>
<td>34</td>
</tr>
<tr>
<td>13</td>
<td>M</td>
<td>10</td>
<td>37</td>
<td>1-2</td>
<td>Repair V.S.D.</td>
<td>49</td>
<td>54</td>
<td>34</td>
<td>34</td>
<td>35</td>
</tr>
<tr>
<td>14</td>
<td>F</td>
<td>36</td>
<td>65</td>
<td>1-7</td>
<td>Repair A.S.D. (secundum)</td>
<td>24</td>
<td>46</td>
<td>22</td>
<td>35</td>
<td>35</td>
</tr>
<tr>
<td>15</td>
<td>F</td>
<td>35</td>
<td>45</td>
<td>1-4</td>
<td>Repair A.S.D. (secundum)</td>
<td>26</td>
<td>55</td>
<td>28</td>
<td>35</td>
<td>36</td>
</tr>
<tr>
<td>16</td>
<td>M</td>
<td>8</td>
<td>25</td>
<td>0-9</td>
<td>Repair V.S.D.</td>
<td>85</td>
<td>60</td>
<td>24</td>
<td>24</td>
<td>27</td>
</tr>
<tr>
<td>17</td>
<td>M</td>
<td>7</td>
<td>21</td>
<td>0-8</td>
<td>Repair A.S.D. (secundum)</td>
<td>24</td>
<td>47</td>
<td>27</td>
<td>35</td>
<td>35</td>
</tr>
<tr>
<td>18</td>
<td>M</td>
<td>34</td>
<td>75</td>
<td>2-0</td>
<td>Repair A.S.D. and I.V.C. draining into L.A.</td>
<td>76</td>
<td>40</td>
<td>32</td>
<td>33</td>
<td>34</td>
</tr>
<tr>
<td>19</td>
<td>M</td>
<td>23</td>
<td>63</td>
<td>1-7</td>
<td>Repair A.S.D. (secundum)</td>
<td>30</td>
<td>48</td>
<td>28</td>
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<td>35</td>
</tr>
<tr>
<td>20</td>
<td>M</td>
<td>8</td>
<td>26</td>
<td>0-9</td>
<td>Repair V.S.D. and pulmonary valvotomy</td>
<td>56</td>
<td>60</td>
<td>23</td>
<td>35</td>
<td>35</td>
</tr>
<tr>
<td>21</td>
<td>F</td>
<td>8</td>
<td>24</td>
<td>1-0</td>
<td>Repair V.S.D.</td>
<td>46</td>
<td>62</td>
<td>28</td>
<td>35</td>
<td>35</td>
</tr>
<tr>
<td>22</td>
<td>M</td>
<td>18</td>
<td>72</td>
<td>1-9</td>
<td>Repair A.S.D. (sinus venosus)</td>
<td>70</td>
<td>42</td>
<td>26</td>
<td>29</td>
<td>31</td>
</tr>
</tbody>
</table>

The patient. The circulation, as judged by central venous and arterial pressures and right atrial filling, was not embarrassed.

Urine production was very low during operation and did not reach a maximum until between 48 and 72 hours afterwards. Thus the average urine flow for the 22 patients was 13.5 ml./kg in the first 24 hours, 33.5 ml./kg. in the second 24 hours, and 52 ml./kg. in the third 24 hours.

The blood loss during and after operation in the first 14 patients is shown in Fig. 1. Out of the 14 patients, six had a loss of less than 1 litre/m² body surface area. In Table II the blood loss in 109 previous cases, with whole blood primes, is compared with all 22 cases.

**TABLE II**

BLOOD LOSS IN 22 CASES WITH HAEMODILUTION COMPARED WITH 109 PREVIOUS CASES WITH WHOLE-BLOOD PRIMES

<table>
<thead>
<tr>
<th>Cases</th>
<th>Loss in Theatre (l/m²)</th>
<th>Post-op. Loss (l/m²)</th>
<th>Total Loss (l/m²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Present</td>
<td>0.67</td>
<td>0.83</td>
<td>1.5</td>
</tr>
<tr>
<td>Previous</td>
<td>0.81</td>
<td>1.29</td>
<td>2.1</td>
</tr>
</tbody>
</table>
Changes in P.C.V. are shown in Fig. 2, which demonstrates the rapid return to normal in the first hour after perfusion. At the end of perfusion, the arterial blood pH was 7.58, Pco₂ 28 mm. Hg, and standard bicarbonate 27 mEq/l. Plasma sodium fell from 137 mEq/l. one hour after perfusion to 120 mEq/l. on the second post-operative day; K⁺ from 4.2 to 3.4 mEq/l. and Cl⁻ from 102 to 91 mEq/l. Urine flow was 1,190 ml. in the first 24 hours, 1,500 ml. in the next 24 hours, and 2,300 ml. in the third 24 hours following perfusion.

Changes in P.C.V. are shown in Fig. 2 with a similar rise immediately following perfusion. At the end of perfusion, the arterial blood pH was 7.57, Pco₂ 15 mm. Hg, and standard bicarbonate 22.5 mEq/l. Plasma Na⁺ fell from 138 mEq/l. one hour post-perfusion to 135 mEq/l. on the second post-operative day; K⁺ from 4.2 to 3.4 mEq/l. and Cl⁻ from 102 to 91 mEq/l. Urine flow was 1,190 ml. in the first 24 hours, 1,500 ml. in the next 24 hours, and 2,300 ml. in the third 24 hours following perfusion.

Two other patients developed post-operative complications. Case 5 required bronchoscopy 24 hours after operation for retention of sputum in the left bronchus. Case 12 had two convulsions on the day after operation. Both patients subsequently made a normal recovery.

**DISCUSSION**

The use of non-blood fluids in the priming of the extracorporeal circulation has many theoretical disadvantages, such as the reduction in oxygen-carrying capacity, with the danger of tissue hypoxia, dilution of the clotting factors, and electrolyte changes. In practice, these do not exist.

In view of her poor pre-operative general condition, and following a moderately prolonged perfusion, an elective tracheostomy was performed at the end of the operation. There were no respiratory complications, and she was discharged on the eighteenth post-operative day.

**CASE 10** A girl aged 14 years, who had had a previous pulmonary valvotomy, underwent closure of a ventricular septal defect and relief of pulmonary infundibular stenosis. Before operation her Hb was 17 g./100 ml., P.C.V. 52%, and plasma Na⁺ 141 mEq/l., K⁺ 4.4 mEq/l., and Cl⁻ 108 mEq/l. Three litres of Ringer-lactate solution (62 ml./kg.) and 60 mEq sodium bicarbonate were used as the priming fluid.

Perfusion lasted 75 minutes, with a partial perfusion of 42 minutes and hypothermia to 27° C. Pharynx (21-5° C. oesophagus). The P.C.V. fell to 22% after 10 minutes' perfusion, then rose slowly to 28% by the end of perfusion. Maximum flow was 2.2 litres per minute at 27° C., and 3 litres of fluid were returned at the end of the perfusion to the patient. Arterial pressure varied from 40 to 45 mm. Hg during total perfusion, and pupil size from 3 to 6 mm. diameter. During a phase of total perfusion, the maximum flow obtained fell to 500 ml. without obvious cause, but improved at once on partial rewarming.

In view of her poor pre-operative general condition, and following a moderately prolonged perfusion, an elective tracheostomy was performed at the end of the operation. There were no respiratory complications, and she was discharged on the eighteenth post-operative day.
weight, while Roe, Swenson, Hepps, and Bruns (1964) showed that a reduction of P.C.V. to 20 to 24% was well tolerated by humans.

The restoration of the patient's red cell mass to normal at the end of perfusion requires the infusion of a large volume of fluid, most of which must leave the circulation rapidly. Hence plasma expanders are not suitable for use as a total prime. Long et al. (1961) used low molecular weight dextran (L.M.W.D.) as a partial priming fluid to overcome the aggregation of red cells, and to increase the capillary perfusion, in the peripheral circulation. However, L.M.W.D. is not only retained in the circulation for at least four hours but also attracts an equal volume of fluid from the tissues. In addition, they found that more than 20 ml./kg. leads to derangement of the clotting mechanism. Thus it is unsuitable in a total prime.

During perfusion, when the circulating blood volume remained near normal limits, the P.C.V. rose slowly. This indicates either a small transfer of fluid from the vascular to the extravascular compartment or an increasing capillary perfusion secondary to the lowered viscosity. When the R.B.C. mass in the patient had been restored to normal, the P.C.V. of the circulating blood rose to pre-perfusion levels within one hour. This indicates a considerable shift of fluid into the extravascular compartment. Moreover, there is evidence that the infused fluid and electrolytes are not fully excreted until the third post-operative day. The urinary output in the 10 cases reported rose to 52 ml./kg. body weight on that day, and the concentration of electrolytes in the urine increased.

The plasma potassium fell from its pre-operative value, and this was maximal on the first post-operative day. Haupt, Myers, Daly, and Birkhead (1964) found a similar fall. Barnard, Saunders, Eales, and Barnard (1966) have reported this to be due to a redistribution of potassium ions between the intra- and extra-cellular body compartments. It was thought likely that the redistribution was the cause of ventricular arrhythmias occurring after perfusion. Prevention of hyperventilation and of sodium bicarbonate excess could minimize this shift, as could the addition of 3% carbon dioxide to the gases in the oxygenator. It is important to prevent alkalosis during perfusion for this reason. During perfusion, high flow rates were easier to attain and maintain, although in one case (case 10) there was difficulty in obtaining adequate venous return.

The arterial pressure was often low during perfusion, and steady levels of 20 mm. Hg were not unusual. The lowering of the blood viscosity must contribute to this and must improve the perfusion during hypothermia when blood viscosity normally increases. Evidence of adequate tissue perfusion at such pressures is provided by a normal pupil size during perfusion, by the early regain of consciousness, and by the absence of any degree of metabolic acidosis.

Difficulty was experienced in obtaining hypothermic cardiac arrest or ventricular fibrillation during perfusion. In case 7, sinus rhythm persisted in spite of an oesophageal temperature of 18.5°C. This observation was also made by Roe et al. (1964).

After operation, the patients usually recovered consciousness in the theatre, and their cooperation in the immediate post-operative period was good. The incidence of respiratory problems was reduced, and there was a noticeable improvement in the patients' colour when breathing air during the early post-operative period as compared with those who had had whole blood primes. It is realized that these impressions are of limited value and more evidence is being collected.

Previous authors have been impressed by the decreased blood loss despite the theoretical disadvantages of diluted clotting factors (Cooley et al., 1962; Roe et al., 1964). The decreased viscosity, in contrast, leads to less damage to the blood elements. The results in this series, though small, show that the blood loss was 30% less than in the earlier cases (Table II). Thus the haemodilution technique may lead to a saving of 3 litres of blood for priming the extracorporeal circuit and also one or two units of transfused blood per average patient.

**SUMMARY**

The use of Ringer-lactate solution as a non-blood prime for the extracorporeal circulation is described. Twenty-two cases in which this technique has been used are described, and the advantages and disadvantages are discussed. Ringer-lactate appears to be a satisfactory non-blood priming solution in dilutions up to 60 ml./kg. body weight, and with a P.C.V. of not less than 20%.

We thank Mr. D. G. Taylor, who operated on all the patients, for permission to publish these results and for his advice and encouragement. We should also like to thank the sisters and nursing staff of the City General Hospital, Royal Infirmary, and Children's Hospital, and Miss Norma Pinkney for secretarial assistance. The diagrams were prepared by Mr. A. S. Foster, medical artist, and the photographic Department of the Royal Infirmary, Sheffield.
ADDENDUM

Since recording these cases a further 18 patients have been operated on using this technique, none of whom had total Ringer-lactate primes. The results support those reported above.

REFERENCES


Ringer-lactate solution as a priming fluid for the disc oxygenator.
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