During the 50 years that have elapsed since Arneth originally studied the behavior of circulating leukocytes, many substances and physical factors have been shown to exert an effect of either depressing or stimulating the total numbers of circulating white cells. Most of the substances have been protein in nature. Thus bacterial products (1), peptone (2), enzymes (3), tissue extracts (4), and pseudoglobulins (5), have all been shown to cause temporary alteration in the peripheral leukocyte picture. Among the physical factors that can similarly influence white cell circulation are cold (6) and ultraviolet light (7). Boyd (8), in 1913, was the first to show that inorganic ions can stimulate an increase in the total white count. While treating insanity with intravenous salt solution, he noted an increase in the polymorphonuclear cells of the blood. Later Bluemel and Lewis (9), in 1924, were able to demonstrate a transitory leukocytosis under the same circumstances. Beard and Beard (10), in 1928, working with rabbits, showed that the intravenous injection of salt caused a transitory leukopenia followed by a leukocytosis. They postulated that the cation Na⁺ might play some role in regulation of white counts.

While working with leukocyte tissue cultures, it was observed (11) that polymorphonuclear neutrophiles exhibited a lengthened survival in hypertonic media whereas lymphocytes showed a corresponding slightly lengthened survival in hypotonic media. Although such studies dealt solely with cell survival, they raised interesting possibilities of correlation with in vivo effects on regulation of circulating leukocytes. The present study was therefore undertaken to determine whether simple alteration in the toxicity of serum would similarly effect leukocyte maturation and release into the peripheral circulation. That is: would experimental hypertonicity induce a selective polymorphonuclear leukocytosis?

METHODS

Subjects: Healthy 10- to 14-kilogram dogs. Hypertonicity was induced, both acutely and chronically, by diverse means. Prior to the start of any 1 experiment, the animals were maintained on a standard diet and fluid intake for 1 week. In experiments which extended over a several-day period, all blood studies were done at the same time of day, under the same circumstances and by the same set of observers, to minimize, as far as possible, the effects of activity, excitement and variation in the diurnal tide on the peripheral leukocyte pattern. Total white counts and smears were made with standard techniques. Red cell counts and/or hematocrits were done throughout each experiment. All variations in absolute leukocyte counts were thus corrected for the factor of hemoconcentration. All variations in tonicity of the extracellular fluid were determined by the technique of freezing point depression on defibrinated blood and are reported in milliosmoles per liter (conversion factor: ΔTᵣ = 1.86 M).

CHRONIC HYPERTONICITY

Chronic hypertonia was attained over a 5- to 10-day period by the following means: (1) Chronic water and food deprivation; (2) chronic water deprivation in the presence of a dry protein diet; (3) chronic water deprivation in the presence of a dry protein diet plus hypertonic salt solution parenterally. As shown by Danowski (12), and others, the diuretic effect of the end products of nitrogen metabolism hastened the onset of chronic hypertonia. Chronic hypertonia was effected 1 time on each of 3 separate dogs, “A,” “C” and “F.”

RESULTS

The average resting (pre-experimental) molarity was 308 milliosmoles. The levels of hypertonicity that were attained varied from 364 to 484 8 milliosmoles. In each instance the hyper-

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Tonicity was accompanied by an increase in the absolute numbers of circulating neutrophiles. The lymphocytes and eosinophiles remained essentially unchanged. The absolute numbers of monocytes decreased. In Figures 1 and 2, the increase in absolute neutrophile counts are plotted against the rise in molarity which occurred in each of the 3 experiments, A-2, C-1, F-1. It will be seen that the molarity of the serum and the absolute neutrophile count bore a linear relation to one another. Under the experimental conditions which existed, the absolute numbers of circulating polymorphonuclear neutrophiles appeared to be a factor of the tonicity of the extracellular fluids.

ACUTE AND CHRONIC HYPERTONIA
EXPERIMENTS A-3, C-1, F-1

0=EXP. A-3 ACUTE HYPERTONIA — ONE I.V. - 100 ML. - 1.6 MOLAR SALT
●=EXP. C-1 CHRONIC HYPERTONIA — 9 DAYS WATER DEPRIVATION
△=EXP. F-1 CHRONIC HYPERTONIA — 8 DAYS WATER DEPRIVATION PLUS DRY PROTEIN DIET

* CORRECTED FOR HEMOCONCENTRATION

Fig. 1.
ACUTE HYPERTONICITY

Acute hypertonicity was produced, in all instances, by the intravenous administration of hypertonic sodium chloride solution. The experiment was performed 6 times on 2 animals, dogs “A” and “G.” In Experiment A-3, 100 ml. of 1.6 molar sodium chloride were infused. In all other experiments, G-2, 3, 4, 6 and 7, 250 ml. of 1.0 molar sodium chloride solution were infused. If sufficient time was allowed for the administration of the infusions (45 to 90 minutes), this amount of salt was exceptionally well tolerated and produced no objective signs of physiologic disturbance. If, however, the rate of infusion was reduced to 1/4 hour or less, profound cerebral irritability, clonic convulsions and death occurred.

RESULTS

Administration of hypertonic salt solution intravenously resulted in acute hypertonia. The molarity of the serum rose rapidly to an average level of 370 milliosmoles and returned to normal in about 7 hours. In all instances, the rise in molarity was followed by a selective increase in the absolute numbers of circulating neutrophiles. The neutrophile rise was statistically significant and of an order of magnitude of 50 per cent or greater. Immediately following the infusion of the hypertonic salt solution, there was a slight decrease in the total numbers of all cell types, both red and white. This was felt to be due to a simple dilution effect from the volume of infused solution and not to be indicative of any specific action on blood cells. Then the rise in absolute numbers of granulocytes began abruptly, around 1/4 to ½ hour after the infusion was completed, and reached its highest level in from 7 to 10 hours. In acute hypertonicity, as in chronic, the increase in absolute neutrophile count bore a linear relationship to the increase in molarity of the extracellular fluids (Exp. A-3, Figure 1). Since, with acute hypertonicity, the rise in molarity usually preceded by an hour or more, the rise
in neutrophiles, the fluctuations in count and
tonicity are best plotted against time. Thus in
Figures 3, 4 and 5, the 24-hour variation in abso-
lute counts following a single infusion of 250 ml.
of hypertonic saline may be seen for Experiments
G-2, 3 and 4. As a control experiment it was
next decided to test the effect of an infusion of a
similar amount of isotonic saline to determine
how much of the polymorphonuclear rise, if any,
could be attributed to a non-specific "washing
out" of cells secondary to an increased rate of
blood flow.
In Experiment G-5 (Figure 6), it will be seen that a transitory rise in neutrophile count did occur after isotonic saline. The rise was slight, however, and not sustained. It was felt that part of this rise might be due to splenic contraction from the stimulus of the intravenous infusion. The animal was therefore splenectomized and the experiments repeated, with both isotonic and hypertonic saline.

In Experiment G-8 (Figure 7), it will be seen that removal of the spleen completely abolished the non-specific rise that occurred in the control injection of isotonic saline. However, in Experiments G-6 and 7 (Figures 8 and 9), it will be seen that removal of the spleen did not affect the rise in count which follows hypertonicity. The rise in neutrophile count, as previously, was
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prompt, selective (unaccompanied by lymphocytosis) and statistically significant.

DISCUSSION

Within the limits of these experiments it is reasonable to ascribe a cause and effect relationship between the milliosmolar strength of the serum and the absolute numbers of circulating neutrophiles. Due to the fact that the leukocyte variations in these experiments were produced by alterations in osmotic pressure rather than through the use of foreign proteins or other substances alien to the blood stream, it is not unreasonable to postulate that variations in milliosmolar strength of the extracellular fluids may play a role in determining the leukocyte pattern in some diverse disease states. For example: (1) The unexplained polymorphonuclear leukocytosis which regularly accompanies the hypertonicity of diabetic coma. (2) The unexplained granulopenia and
relative lymphocytosis which regularly accompanies the hypotonicity of Addison's disease. Studies are currently being carried on to investigate these relationships.

The mode of action of the hypertonic polymorphonuclear leukocytosis produced in these experiments is unknown. Its prompt occurrence in acute hypertonicity makes it obvious that it is not due to the effect which originally stimulated these investigations, namely: The lengthened survival time of neutrophiles in hypertonic media (11). The recent work of Lawrence (13), who places the in vivo survival time of leukocytes at 16 hours, would similarly tend to make such an effect improbable. The 2 most likely mechanisms would seem to be either a release of preformed cells from some theoretical "store house" in the body or an accelerated maturation and release of immature cells from the bone marrow. If the first mechanism obtains, it would be difficult to visualize the absence of a simultaneous rise in all leukocyte types (lymphocytes, monocytes,
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**Dog-G Exp. B Control**

Effect of single injection of 250 ml. isotonic salt solution i.v. on splenectomized animal.

**Summary**

Hypertonicity of the serum of dogs was produced both acutely and chronically. In all in-

Eosinophiles) and the persistence of the rise in the case of the chronically induced hypertonicity. If the second mechanism obtains, one would expect rather large numbers of juvenile neutrophiles to appear in the peripheral circulation. Although this phenomenon was looked for, and sometimes found, its appearance was too irregular to lend weight to this possibility. On the basis of these experiments, no explanation of the mechanism can be offered. Further investigation into the mode of action of hypertonic polymorphonuclear leukocytosis should include serial bone marrow observations.
stances it was accompanied by relative and absolute increases in the numbers of circulating neutrophiles. The absolute numbers of lymphocytes remained essentially unchanged with increases in molarity. Monocytes, when present in numbers to be statistically significant, showed a slight decrease with increases in molarity.

CONCLUSION

Under the conditions of these experiments the increases in absolute numbers of circulating neutrophiles were directly proportional to the increases in the milliosmolar strength of the extracellular fluids.
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DOG-G EXP. 7 ACUTE HYPERTONIA EFFECT OF SINGLE INJECTION OF 250 ML.-1 M. SALT SOLUTION I.V. ON SPLENECTOMIZED ANIMAL

FIG. 9.
BIBLIOGRAPHY