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Authors

Schooley, John C
Mahlmann, Lynn J

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STIMULATION OF ERYTHROPOIESIS IN PLETHORIC MICE BY PROSTAGLANDINS
AND ITS INHIBITION BY ANTI-ERYTHROPOIETIN*

By

John C. Schooley and Lynn J. Mahlmann

Lawrence Radiation Laboratory, Donner Laboratory,
University of California, Berkeley, California 94720

Running Title: Prostaglandins and erythropoiesis

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Prostaglandins, particularly of the E series, are potent vasodilators of most vascular beds, and some attention has been devoted to their possible role in the control of renal blood flow (1-3). Considerable evidence indicates that the primary regulation of plasma erythropoietin resides in the kidney (4). It appeared possible that prostaglandins might redistribute renal blood flow in plethoric mice, and thus stimulate erythropoiesis. The present studies were undertaken to determine whether erythropoiesis was stimulated by prostaglandins in the rodent, and whether the stimulation was erythropoietin-dependent.

Materials and Methods

Female LAF₁/Jax mice weighing about 25 g were used. Mice were made plethoric by exposure to increasing amounts of carbon monoxide as described by Fogh (5). The mice were used 7 days after removal from the chamber when erythropoiesis was maximally suppressed.

Prostaglandins (PG) E₁, E₂, and F_{2α} were obtained from Dr. John E. Pike of the Upjohn Company. They were dissolved in one ml of ethyl alcohol and diluted with saline to 100 ml to give a final concentration of 100 µg/ml. Mice were injected sc with 1 µg/g body weight. Some mice received the PGE₁, and PGE₂ on two consecutive days.

Anti-erythropoietin immune serum obtained from rabbits immunized with human urinary erythropoietin was injected iv in a volume of 0.1 ml immediately before each PGE₁ or PGE₂ injection. One ml of this particular immune serum can neutralize the biological activity of 25 I.R.P. units of human erythropoietin or about 2.5 units of sheep or mouse erythropoietin. The method of preparation and properties of anti-erythropoietin have been summarized elsewhere (6). Control groups received similar injections of normal rabbit serum.

Groups of plethoric mice were exposed to a simulated altitude of 22,000 ft

(321 torr) for 6 hr; separate groups were injected with PGE₁, PGE₂, or PGF_{2α} immediately before the hypoxic exposure, and one group received saline.

Fifty-six hours after the first prostaglandin injection the plethoric mice were injected iv with 0.5 μCi of ⁵⁹Fe as the citrate. All mice were bled 72 hr after the ⁵⁹Fe injection, and the radioactivity in 0.5 ml of blood was measured. The results are expressed as the percent of the injected ⁵⁹Fe in the calculated blood volume. The blood volume of these plethoric mice was assumed to be 7% of the body weight. The average hematocrit of all groups was 64.1 ± 1.1% at the end of the assay. Each experimental group contained 6-8 assay mice.

Results

The data shown in Table I indicate that a single sc injection of PGE₁ and PGE₂, at a dose of 1 μg/g body weight, significantly stimulates erythropoiesis in plethoric mice, as measured by ⁵⁹Fe incorporation. PGF_{2α} at the same dosage did not stimulate erythropoiesis.

The data in Table I also show that prostaglandins of the E series significantly (P < 0.001) potentiate the erythropoietic response of plethoric mice exposed to a 6 hr hypoxic exposure, increasing the ⁵⁹Fe incorporation about two-fold. Significant potentiation was not observed with PGF_{2α} (P > 0.1).

The data presented in Table II indicate that the erythropoietic response elicited by PGE₁ and PGE₂ injected sc on two consecutive days is completely abolished when anti-erythropoietin is injected before each PG injection.

Discussion

These experiments clearly indicate that prostaglandins of the E series significantly stimulate erythropoiesis in the plethoric mouse as measured by the 72-hr ⁵⁹Fe incorporation into red blood cells. It is equally clear that this stimulation is erythropoietin-dependent, since it does not occur in

anti-erythropoietin injected plethoric mice.

It is tempting to speculate that, among the wide variety of physiological and pharmacological actions of prostaglandins (1-3), their role as vasodilators of vascular beds is involved in the observed erythropoietin-dependent stimulation of erythropoiesis. Lee (7) has shown that the renal blood flow of dogs is increased after PGE_1 and PGE_2 , but not $PGF_{2\alpha}$. Possibly similar changes occur in the mice used in these experiments after PGE_1 and PGE_2 injection, causing a redistribution of renal blood flow with a decreased O_2 tension in some appropriate tissue, resulting ultimately in an increased erythropoietin production. The failure of $PGF_{2\alpha}$ to stimulate erythropoiesis may relate to its failure to alter renal blood flow. However, only further experiments can clarify the mechanism of prostaglandin stimulation of erythropoiesis.

The ubiquitous distribution of prostaglandins in almost all mammalian tissues considered, along with the results of the present experiments, suggests that some caution is required before concluding that the erythropoietic stimulation observed after injection of various tissue extracts into plethoric mice is the direct result of the presence of erythropoietin in the extract.

Summary

Prostaglandins of the E series stimulated erythropoiesis in the plethoric mouse, whereas prostaglandin $F_{2\alpha}$ was inactive. The erythropoietic stimulation was erythropoietin-dependent, since it did not occur in anti-erythropoietin injected mice.

Table I

Erythropoietic response of plethoric LAF₁ mice to prostaglandins
with or without a brief hypoxic exposure

	<u>72-hr ⁵⁹Fe incorporation</u>
Saline	0.60 ± 0.02*
PGE ₁	2.77 ± 0.41
PGE ₂	3.84 ± 0.57
PGF _{2α}	0.73 ± 0.05
6 hrs. hypoxia**	12.7 ± 2.4
6 hrs. hypoxia + PGE ₁	23.6 ± 1.4
6 hrs. hypoxia + PGE ₂	26.7 ± 0.77
6 hrs. hypoxia + PGF _{2α}	19.5 ± 3.1

* Standard error of the mean.

** Simulated altitude of 22,000 ft (321 torr). Prostaglandins given
once sc, 1μg/g body weight.

Table II

Erythropoietic response of plethoric IAF₁ mice receiving
prostaglandins and anti-erythropoietin

	72-hr ⁵⁹ Fe incorporation*
Saline	0.91 ± 0.10
PGE ₁ + Normal rabbit serum	5.71 ± 0.36
PGE ₁ + Rabbit anti-erythropoietin	0.62 ± 0.06
PGE ₂ + Normal rabbit serum	6.80 ± 1.1
PGE ₂ + Rabbit anti-erythropoietin	0.66 ± 0.07

* Standard error of the mean. Prostaglandins given sc on two consecutive days 1μg/g body weight. 0.1 ml of immune sera after each PG injection.

References

1. Bergström, S., Carlson, L. A., and Weeks, J. R., *Pharmacol. Revs.* 20, 1 (1968).
2. Horton, E. W., *Physiol. Rev.* 49, 122 (1969).
3. Ramwell, P. W. and Shaw, J. E., *Rec. Prog. Horm. Res.* 26, 139 (1970).
4. Krantz, S. B. and Jacobsen, L. O., "Erythropoietin and the Regulation of Erythropoiesis," 330 pp. Univ. Chicago Press, Chicago (1970).
5. Fogh, J., *Scandinav. J. Clin. Lab. Invest.* 18, 33 (1966).
6. Schooley, J. C. and Garcia, J. F., *Blood* 25, 204 (1965).
7. Lee, J. B., in "Prostaglandin Symposium of the Worcester Foundation for Experimental Biology" (P. W. Ramwell and J. E. Shaw, eds.), p. 131 John Wiley & Sons, New York (1968).