Proceedings of the Iowa Academy of Science

Volume 78 | Number 1-2

Article 9

1971

Effect of Salicylic Acid on the Early Development of the Chick Limb-Bud

Suleiman A. Suleiman *Iowa State University*

John R. Baker Iowa State University

Copyright ©1971 Iowa Academy of Science, Inc. Follow this and additional works at: https://scholarworks.uni.edu/pias

Recommended Citation

Suleiman, Suleiman A. and Baker, John R. (1971) "Effect of Salicylic Acid on the Early Development of the Chick Limb-Bud," *Proceedings of the Iowa Academy of Science, 78(1-2),* 20-23. Available at: https://scholarworks.uni.edu/pias/vol78/iss1/9

This Research is brought to you for free and open access by the Iowa Academy of Science at UNI ScholarWorks. It has been accepted for inclusion in Proceedings of the Iowa Academy of Science by an authorized editor of UNI ScholarWorks. For more information, please contact scholarworks@uni.edu.

Proc. Iowa Acad. Sci. 78 (1971)

Effect of Salicylic Acid on the Early Development of the Chick Limb-Bud

SULEIMAN A. SULEIMAN¹ and JOHN R. BAKER²

SULEIMAN A. SULEIMAN and JOHN R. BAKER. Effect of Salicylic Acid on the Early Development of the Chick Limb-Bud. Proc. Iowa Acad. Sci., 78(1):20-23, 1971.

SYNOPSIS. Transplanted hind limb-buds of chick embryos were treated with salicylic acid. Treatment with this chemical caused reduction in length, weight and volume of the developing limbbuds. A side effect observed was an apparent loosening of the and examined with regard to length, weight, and volume.

Mitochondria are organelles found in all aerobic organisms in which the oxidation of pyruvic acid to carbon dioxide and water takes place.

Several workers have isolated mitochondria to study their properties and find their role in embryogenesis. Carey and Greville (1959a,b) found that the mitochondria obtained from embryonic preparations appeared to be more fragile than their adult counter parts. McAlpino (1960) reported a marked increase in phosphate activity (which is related to mitochondria) in the ectodermal tips of the developing limbbuds of rat embryos. These observations and others (Bosund, 1957; Charnock, and Opit 1962a) show an essential mitochondrial activity during early development.

Salicylates are thought to be inhibitors of mitochondrial activity, which may act by disrupting or uncoupling the link between oxidation and phosphorylation. Brody (1956) was the first to show that salicylates in a concentration above 3 mg/100 ml. can decrease the P:O (phosphorylation: oxidation) ratio of rat liver and kidney mitochondria oxidizing a variety of substrates. The actual site of salicylate activity has not yet been found; but Charnock and Opit (1962b) have suggested that the locus of the action of salicylate in uncoupling oxidative phosphorylation is at the level of the mitochondrial membrane. It was therefore considered desirable to observe the effect of Salicylic acid on the early development of the chick limb-buds and to find if Salicylic acid has another effect in addition to its suggested uncoupling factor.

METHODS AND MATERIALS

The hind limb-buds of white leghorn chick embryos at stage 19 of Hamilton and Hamburger, were extirpated and transplanted singly onto the chorioallantoic (CA) membrane of host embryos of stage 35 of Hamilton and Hamburger. Immediately after transplantation, 0.08 ml. of salicylic acid were dropped on top of the transplanted buds with a microdropper. There were seven groups of transplants corresponding to seven concentrations. The host eggs were again incubated at 38° C. The transplanted limb-buds were recovered at intervals of 6 hours, 1, 2, 3, 4, and 5 days after operation basal membrane. The above observations are discussed in relation to the different hypotheses of the ectoderm-mesoderm interaction. The physiological role of the basal membrane, as well as the possible implications of the basal findings to the question of limb development and morphogenesis, are also discussed. INDEX DESCRIPTORS: Salicyclic acid, chick limb bud, ectoderm-

mesodern interaction, limb bud growth.

Some of them were then fixed in 4 per cent neutral formalin and others in Helley's fluid, washed in tap water, dehydrated, and embedded in paraffin. The sections were cut at 4 u by AP-121 Microtome, and stained either by Novelli's method for demonstration of mitochondria or with eosine and haematoxylin (E and H). Controls were done with each group of experimental concentrations in which 0.08 ml. of Pannet-Compton solution were added to the transplant.

For thin sectioning, limb-buds were fixed in chilled gluteraldehyde (4°C) for 4 hours, and post fixed in osmium tetroxide (in phosphate buffer) for 45 minutes. After dehydration with graded alcohols (35, 70, 90 per cent and three changes of absolute alcohol) the tissues were transfered to the Araldite mixture, the Araldite was changed twice or thrice during this time. The sections ($\frac{1}{2}$ - $\frac{3}{4}$ µ thick) were cut by glass knives on LKB ultra-microtome, and were stained with acid fuschin.

OBSERVATIONS

The group of limb-buds treated with 250 μ gm/ml and 200 μ gm/ml concentration showed a pronounced growth inhibition during the whole time of transplantation, and especially during the first 4 days of transplantation, compared to the controls (Figure 1). The groups of limb-buds treated with the concentrations of 170, 150, and 135 μ gm/ml also showed growth inhibition but to a lesser degree and in the period of the first 3 days of transplantation (Figure 2). The groups of limb-buds treated with the concentration streated with the concentration of 125 and 100 μ gm/ml did not show any significant growth inhibition; (Figure 3) and the growth of the group treated with 100 μ gm/ml was very similar to the controls. The only difference between the groups of limb-buds treated with the last two concentrations and the control was less growth in the first day (Figure 3).

In all groups of transplanted limb-buds (except for those treated with the first two concentrations) the growth (length, weight, and volume) tends to become normal after the third day of treatment. In the groups treated with the last two concentrations the growth is very close to the normal. In all the transplanted limb-buds there was no significant difference in growth during the first 6 hours of transplantation compared to the controls. After this the growth behavior differed according to the concentration.

1

¹Suleiman A. Suleiman, 253 Science Building, Iowa State University, Ames, Iowa 50010

²John R. Baker, Ph.D., Assistant Professor of Zoology, 351 Science Building, Iowa State University, Ames, Iowa 50010

LIMB BUD DEVELOPMENT



Fig. 1. Growth of transplanted limb-buds treated with $250\mu gm/$ ml of salicylic acid.

In the untreated limb-buds (Figure 4) the basal side of the ectoderm is sharply demarcated from the mesoderm. It was observed that in Salicylic acid treatment with the higher concentrations (250 μ gm/ml and 200 μ gm/ml) the basal membrane looked loose under the light microscope, and it was easy to separate an almost intact ectoderm from the examined transplanted limb-buds in the period of the third day of transplantation.

In the case of the lower concentrations the loosening was less significant and it was more difficult to separate the ectoderm (Figure 6). In the groups receiving 150 μ gm/ml and 100 μ gm/ml concentrations there was no loosening except after six hours treatment. Whether any change has occured in the chemical constitution of the basal membrane is not known.

DISCUSSION

During the past twenty years, much experimental work has been directed to the study of ectoderm-mesoderm interaction in limb morphogenesis in avian embryos. It has been debated whether the ectoderm or the mesoderm of the limb-bud is the site of developmental factors. Balinsky (1956) suggested that both mesoderm and ectoderm take an active part in the development of an induced as well as normal limb.

Recently two different hypotheses have been formulated to account for the development of limbs in tetrapod vertebrates. In both hypotheses the mesoderm is claimed to be the essential site for limb development. The area of disagreement is in the role played by limb-bud ectoderm, particularly the apical ridge. An increasing body of evidence has accumulated which



Fig. 2. Growth of transplanted limb-bud treated with 170 $\mu gm/$ ml of salicylic acid.



Fig. 3. Growth of transplanted limb-buds treated with 100 $\mu gm/$ ml of salicylic acid.

Proc. Iowa Acad. Sci. 78 (1971)



Fig. 4. Portion of sagittal section of untreated limb-bud recovered after one day; section at $1/2~\mu$ and stained with acid fuschin. X 960.



Fig. 6. Portion of sagittal section of limb-bud treated with 170 μ gm/ml of salicyclic acid recovered after one day; section at 3 μ and stained by Novelli's method. X 1110.



In various publications the term basal membrane is applied to various structures, in this paper the basal membrane is defined as the zone present at the dermo-epidermal junction. In electron microscopy the term is usually applied to a continuous membrane covering the basal membrane cells (Sjostrand, 1953; Jurand, 1965).

It has been suggested that the basal membrane composed mainly of tropocollagen fine filaments which are either embedded in an amorphous matrix of the same density (probably mucopoly-saccharides), or are very closely compacted so that the individual fiber can not be resolved (Fawcett, 1966).

Although the exact function of the basal membrane is still uncertain, it is believed to be a diffusion barrier (Balinsky, 1956, 1957). Caeser and Edward (1957) suggested a protection function from too rapid ion concentration changes.

It is not known if salicylic acid acts on the tropocollagen in the basal membrane, but it interferes with the metabolism of collagen (Bellamy, 1963). It affects the biosynthesis of mucopolysaccharides in connective tissues (Bostrum, 1955, 1963). The loosening of this membrane observed in our experiments may be attributed to this effect, or it might suggest an effect of salicylic acid on other components not yet known. The inhibition of growth by salicylic acid disappears especially in the groups treated with 100-170 μ gm/ml concentrations (Figures 1, 2, 3). This disappearance may be explained by diffusion of salicylic acid into the surrounding tissue. It is also likely that it affects an interaction between the ectoderm and mesoderm through this membrane. ing tissue. It is also likely that it affects an interaction be-Its diffusion and reduction of its effectiveness allows a reformation of an "interacting substance" in this membrane, so a normal or very close to normal growth is maintained again.

More biochemical studies must be done to determine the precise make-up of the basal membrane, and the exact mode of action of salicylic acid on this membrane.



Fig. 5. Portion of sagittal section of limb-bud treated with 250 μ gm/ml of salicylic acid recovered after two days; sectioned at 3 μ and stained by Novelli's method. Note the loosening of the basal membrane. X 1170.

indicates an active participation of ectoderm in limb development (Zwilling, 1961; Amprino, 1962; DeHaan and Ebert, 1964) especially the thickened portion which is morphologically prominent in birds. According to our observation of

LIMB BUD DEVELOPMENT

LITERATURE CITED

- AMPRINO, R. 1962-1963. Aspetti della Morphogenesi della Estremita nei Vertebrati. Mon. Zool. Ital. Suppl. 70-71:7-130.
- BALINSKY, B. I. 1956. A new theory of limb induction. Proc. Acad. Sci., U.S.A. 42:781-785.

_____. 1957. New method on the mode of action of the limb inductor. J. Exp. Zool. 134:239-274.

BELLAMY, A., A. K. HUGGINS, & M. J. H. SMITH. 1963. Salicylate and glutamate metabolism. J. Pharm. Pharmacol. 15:559-560.

BOSTRUM, H. and B. MAANSSON. 1955. The action of salicylates and related compounds on the sulphate exchange of chondroitin sulphuric acid. J. Pharm. Pharmacol. 7:185-190.

A. MORETT & M. WHITEHOUSE. 1963. Studies on the biochemistry of heart valves, 1. on the biosynthesis of mucopolysaccherides in bovine heart valves. *Biohem. Biophys. Acta* 74: 213-221.

BOSUND, I. 1957. The effects of salicylic acid, benzoic acid and some of their derivatives on oxidation phosphorylation. Acta, Chem. Scand. 11:541-544.

BRODY, T. M. 1956. Action of sodium salicylate and related compounds on tissue metabolism in vitro. J. Pharm. Expt. Therap. 117:39-51.

- CAREY, H. H. & C. D. GREVILLE. 1959a. Mitrochondria from embryonic tissues of the chick. 1. preparation, characterization, and some enzymic properties. *Biochem. J.* 71:159-166.
- & ______. 1959b. Mitochondria from embryonic tissues of the chick II. Metabolic activities. *Biochem. J.* 71:166-176.

CHARNOCK, J. S. & L. J. OPIT. 1962a. Effect of salicylates on ATP activity of rat liver mitochondria. *Biochem. J.* 83:596-602.

& ______ & _____. 1962b. An evaluation of the effect of salicylate on oxidative phosphorylation in rat liver mitochondria. *Biochem. J.* 83:602-606.

DEHANN, R. L. & J. D. EBERT. 1964. Morphogenesis. Ann. Rev. Physiol. 26:15-46.

- FAWCETT, D. W. 1966. An atlas of fine structure. The Cell. Philadelphia and London. Saunders.
- JURAND, A. 1965. Ultrastructural aspects of early development of the fore-limb buds in the chick and the mouse. *Proc. R. Soc. B.* 162:387-405.
- MCALFINO, R. J. 1955. Alkaline phosphatase in the developing thyroid, parathyroid and thymus of the albino rat. Am. J. of Ant. 96:191-227.
- SJOSTRAND, F. S. 1953a. Ultrastructure organization of retinal rods and cones. J. Applied Physics 24:117.
- ZWILLLING, E. 1961. Limb morphogenesis. 1. Advances in morphogenesis. New York Academic Press: 301-330.

23

CAESER, R. & G. A. EDWARD. 1957. The physiologic significance of the basement membrane. *Anat. Rec.* 128: 530.