Prolactin release induced by suckling in lactating rats after L-DOPA treatment

R. P. Deis and J. Prilusky

Instituto de Investigacion Médica, Mercedes y Martin Ferreyra, Casilla de Correo 389, Cordoba, Argentina

In a previous paper (Prilusky & Deis, 1975) a blocking effect of L-DOPA on milk ejection and on prolactin release was demonstrated in lactating rats. The inhibitory effect of brain catecholamines on prolactin release is well established and it is probably mediated through dopamine (MacLeod, 1969; Donoso, Bishop, Fawcett, Krulich & McCann, 1971). Since a catecholaminergic system in the central nervous system may regulate prolactin release in lactating rats, we investigated the effect of the suckling stimulus on prolactin release in rats treated with L-DOPA.

White primiparous lactating rats weighing about 270 g were used. Each litter was reduced to six young 4 days after birth. On the 10th day of lactation, the litter was separated from the mother in the early morning for 9 hr. The young were returned at 15.00 hours and allowed to suck for 30 min. The young were weighed to the nearest 0.1 g immediately before and after the suckling period to determine the amount of milk ejected by the mother. The procedure was repeated on 2 consecutive days and on the 3rd day blood samples were obtained, one from each rat, by heart puncture without anaesthesia at various times after the end of the suckling period. This simple method takes only a few seconds and does not disturb the rats (Vermouth & Deis, 1974).

Rats in Group 1 were injected i.p. with L,3-4-dihydroxyphenylalanine (L-DOPA: Regis Chemicals; 10 mg/100 g body wt) suspended in physiological saline 30 min before the suckling period. The serum prolactin levels at 0, 30, 60, 90, 120, 150, 180 and 210 min after the suckling period were measured by radioimmunoassay with two dose levels of serum (Niswender, Chen, Midgley, Meites & Ellis, 1969) and reagents supplied by the NIAMDD. All the serum samples were assessed in a single radioimmunoassay to eliminate between-assay variation. Student’s t test was used to assess the level of significance.

The normal release of prolactin induced by suckling was prevented by L-DOPA during the first 90 min after suckling (Text-fig. 1). A peak in serum prolactin concentration was obtained at 120 min (P<0.001). At 150 min serum prolactin was still significantly higher than the pre-suckling values of control rats (P<0.005), but a progressive decrease occurred at 180 and 210 min after the suckling period. The mean values at 180 and 210 min were not significantly different from the control pre-suckling level and the levels obtained in the first 90 min after suckling in the treated rats.

In order to see if the peak serum prolactin level obtained 120 and 150 min after the suckling period in rats treated with L-DOPA was due to the suckling stimulus and was not a secondary effect of L-DOPA at the central nervous system, rats in Group 2 were similarly treated with L-DOPA but the suckling stimulus was omitted. The serum prolactin concentration was estimated at 90, 180, 210 and 270 min after the L-DOPA injection, corresponding to the 30, 120, 150 and 210 min times for the Group 1 rats. At none of the times studied was the serum prolactin level significantly different from the mean value in control rats before suckling and no peaks were observed.

Four rats in Group 3 were treated with L-DOPA and the effect of the suckling stimulus at the central nervous system was prevented by i.p. injection of sodium pentobarbitone (Nembutal: Abbott; 2 mg/100 g body wt) 30 min before the suckling period. The mean (±S.E.M.) serum prolactin level 120 min after suckling (17.8±5.6 ng/ml) was significantly less than the values obtained in Group 1 (59.3±112.6 ng/ml; P<0.001) and Group 2 (47.8±21.6 ng/ml; P<0.01) rats.

As shown in Text-fig. 1, the inhibitory effect of L-DOPA on the secretion of prolactin after suckling lasted for about 90 min and thereafter a significant increase in serum prolactin concentration occurred. That the peak of serum prolactin was not due to a secondary effect of L-DOPA at the hypothalamic or pituitary level is shown by the results when the suckling stimulus was absent or its
Text-fig. 1. Mean (±S.E.M.) serum prolactin concentrations in untreated lactating rats (Δ) before and after a 30-min suckling period (S) and in rats treated with L-DOPA and with (○) or without (●) a suckling stimulus. The numbers of rats in each group are shown in parentheses.

The present and previous results (Prilusky & Deis, 1975) indicate that monoamines, probably dopamine, may have an important function in controlling prolactin release in lactating rats.

We thank Mr E. Molina for the care of the animals and technical assistance. The work was supported by a grant from the Consejo Nacional de Investigaciones Científicas y Técnicas of Argentina, of which R.P.D. is a career scientist and J.P. a post-doctoral fellow. We are grateful to Labora-
torios Massone, Argentina, for the gift of L-DOPA. The NIAMDD rat pituitary hormone distribution program kindly supplied the radioimmunoassay kit for prolactin.

References


Received 4 July 1975