

Peaks and valleys of prolactin-related gene expression during pigeon parental care stages

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Abstract

The goal of this research was to provide the most comprehensive gene expression profile of gene expression activity in the HPG axis of male and female rock doves to provide a deeper undertand of how the reproductive axis response to typical behavioral transitions associated with parental care. Here we used RNA sequencing to measure gene activity at 8 stages of the parental care cyle from nest building, to egg incubation, and through nestling care. Non-parental groups were added as controls, but gene expressions differences were assesed between and across all time points. The three tissue have unique signatures, but within each tissue we find minial sex differences. The pitutary samples displacy the msot plasticity in gene expression, with many changes in gene expression mirroring the typical rise and fall of circulating prolactin that peaks when chicks hatch. Our analysis provides new insight into how suites of genes respond in concert to the demands of offspring care. This data can be used to develop and test hypotheses about the mechanism regulating parental care behavior.

Author summary

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Introduction

Understanding the mechanisms underlying parental care are critical to circumventing issues with parent-newborn bonding as well, where ultimate explanations are obvious, but specific mechanisms remain elusive. The rock dove (*Columba livia*) is an ideal system to characterize changes in genetic expression during parental care transitions because: 1) ample genomic resources are available (Gillespie et al. 2013), including a complete annotated genome assembly [1] and methodology concerning reproductive physiology and behavior (Dong et al. 2012); and 2) rock doves are prolific, year-round breeders that thrive in captivity, making observation, manipulation, and sampling highly feasible year-round.

Rock doves are socially monogamous and offer bi-parental care, making inter- and intra-sexual comparisons possible. Birds offer two important behavioral transition points into parental care: the incubation of eggs and the caring for chicks. This produces two unique opportunities to study how the brain transitions into two different suites of parental care behaviors. Additionally, rock doves exhibit a parental care

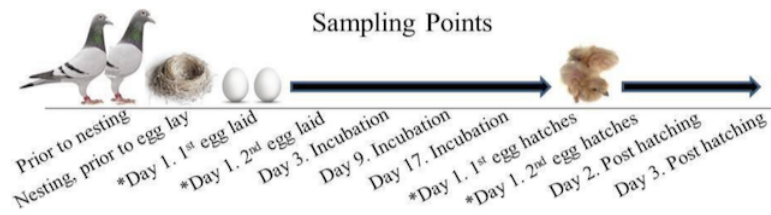
strategy analogous to mammals in that they, too, ‘lactate’ to feed their young (Gillespie et al. 2011, 2012). This lactation, unlike simple regurgitation of food, consists of the production and sloughing off of skin cells inside the crop sac of females and males, creating a protein-rich milk-like substance on which they rear their chicks. Many functional similarities between rock dove and mammalian lactation exist concerning the mediation of this event by the hormone prolactin (Dumont 1965). Additionally, like mammalian milk, rock dove milk delivers essential immunoglobulins and nutritional benefits to young, aiding in their immune function and development of microbiota [2]. Thus, because rock doves incubate eggs and exhibit mammalian-like mediation and function of lactation for young, they have the potential to serve as a powerful theoretical bridge to understand the neurobiology of both avian and mammalian transitions into parental care. Our working hypothesis is that distinct changes in transcription occur in the brain at the anticipation of, during, and in response to two different types of parental care: incubation behavior and hatchling care.

Materials and Methods

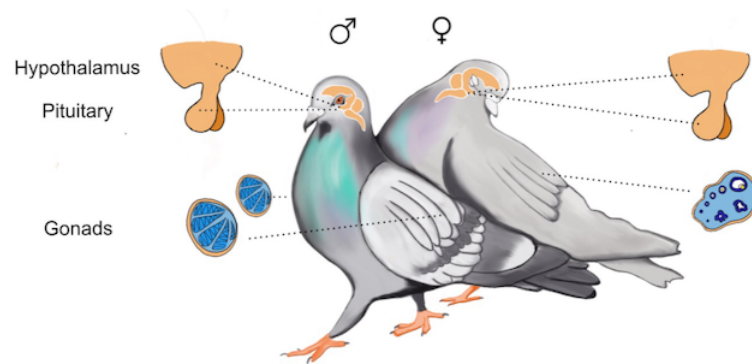
Rock dove housing

Sexually mature male and female rock doves were housed together in rooftop aviaries on the Barnard Campus. Natural male-female pairings permitted, and small observation windows behind each nest box (64 boxes total) facilitate observations nest building, egg laying, chick hatching, and nesting care every morning from 0900-1100 (correct time?). On specific collection days (Fig 1A, needs to be updated), truck blood, the hypothalamus, the pituitary, and the gonads were collected following common protocols used to sample tissue from birds (Fig 1B). Tissue collection, RNA isolation, cDNA library preparation, sequencing, quality control, and read processing were done as described in Calisi et al. 2018, Lang et al 2019, and Austin et al 2019.

A



B



We used EdgeR and DESeq2 to calculate statistical differences in gene expression between each transition. All subsequent analyses were conducted using variance stabilized data from the DESeq2 model.

Results

Thousands of genes were found to be differentially expressed between the non-parental control group and the building group (Table 1, Fig. 1). Within the time course of parental care, fewer than 1000 genes change their expression over the course of a few days, with the most pronounced difference in total number of changes occurring in the female pituitary between incubation days 9 and 17 (Table 1, Fig. 1).

Method	Contrast	Hyp F	Hyp M	Pit F	Pit M	Gon F	Gon M
EdgeR	Cont - Bldg	14674	117	14593	144	14859	4
EdgeR	Bldg - Lay	14742	0	14593	0	14836	0
EdgeR	Lay - Inc3	0	0	0	0	19	0
EdgeR	Inc3 - Inc9	0	1	0	3	2	0
EdgeR	Inc9 - Inc 17	0	2	57	37	9	0
EdgeR	Inc17 - Hatch	0	0	3	1	3	0
EdgeR	Hatch - Nest5	1	0	3	0	0	0
EdgeR	Nest5 - Nest9	0	0	0	0	24	0
DESeq2	Cont - Bldg	3452	4499	3950	4680	4597	2135
DESeq2	Bldg - Lay	0	0	101	0	104	1
DESeq2	Lay - Inc3	0	0	18	1	440	1
DESeq2	Inc3 - Inc9	1	0	0	1	8	0
DESeq2	Inc9 - Inc 17	1	0	640	186	1	0
DESeq2	Inc17 - Hatch	2	0	3	18	1	0
DESeq2	Hatch - Nest5	166	0	340	3	1	0
DESeq2	Nest5 - Nest9	0	0	1	23	19	0

Table 1. Comparison of total DEGs determined by edgeR and DESeq2

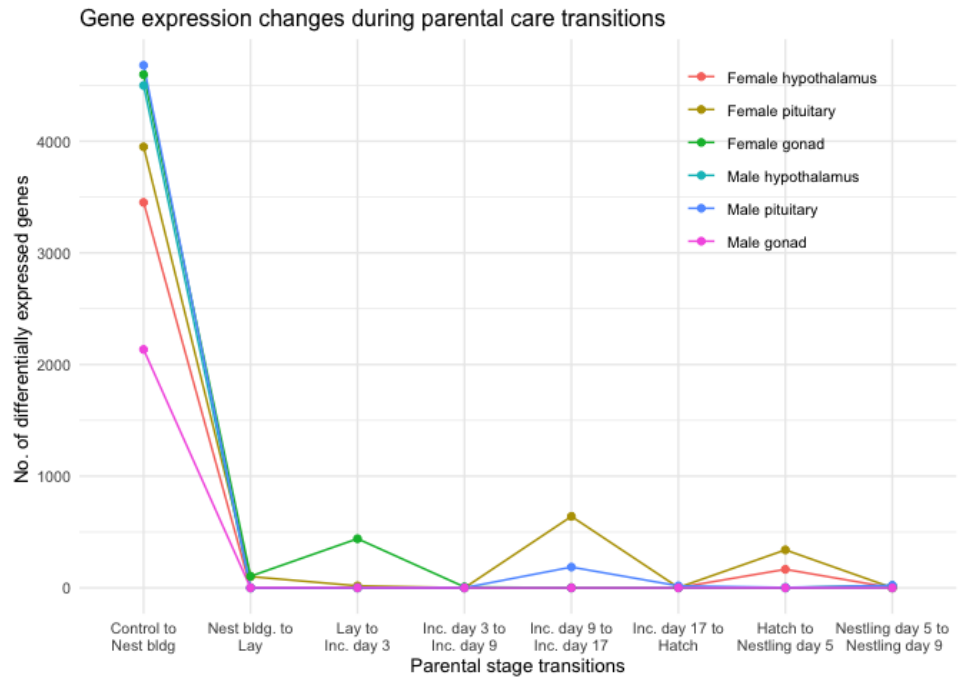


Fig 1. The magnitude of gene expression changes between each parental transition. Using DESeq 2, we identified 4-5K genes are differentially expressed between control birds and their nest building conspecifics in all tissues except the male gonad. 500 - 1000 genes are differentially expressed in the female pituitary from mid-late incubation as well as in the female hypothalamus and pituitary from hatch to nestingly

Fig 2: Ability to use gene expression to distinguish groups

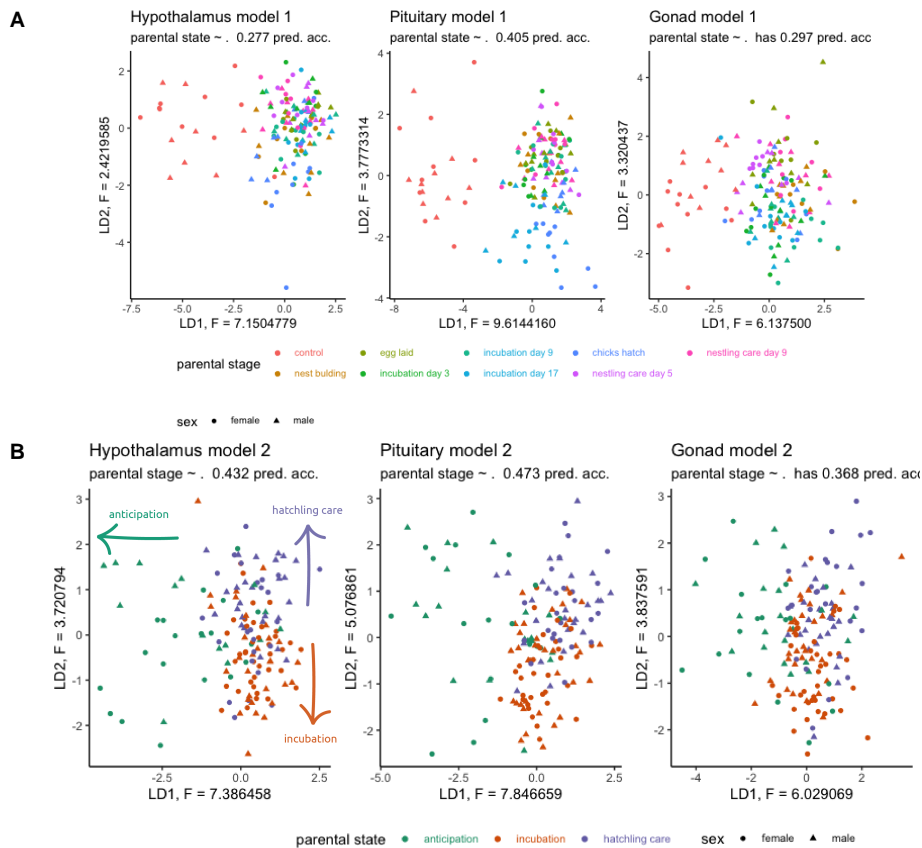


Fig 3: Prolactin expression in PIT (but not HYP or GON)
follows same pattern as in blood

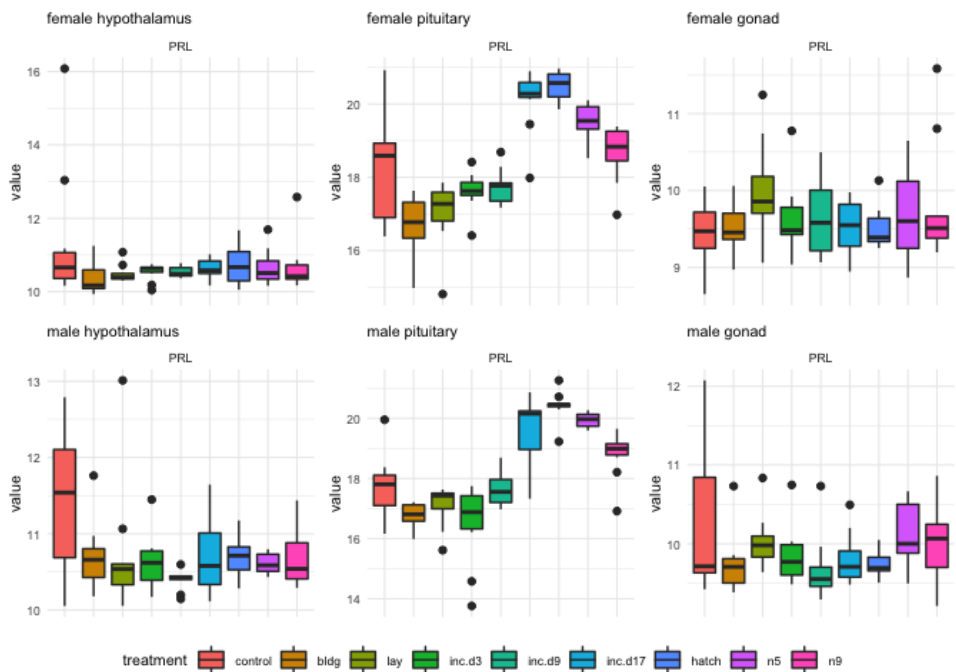
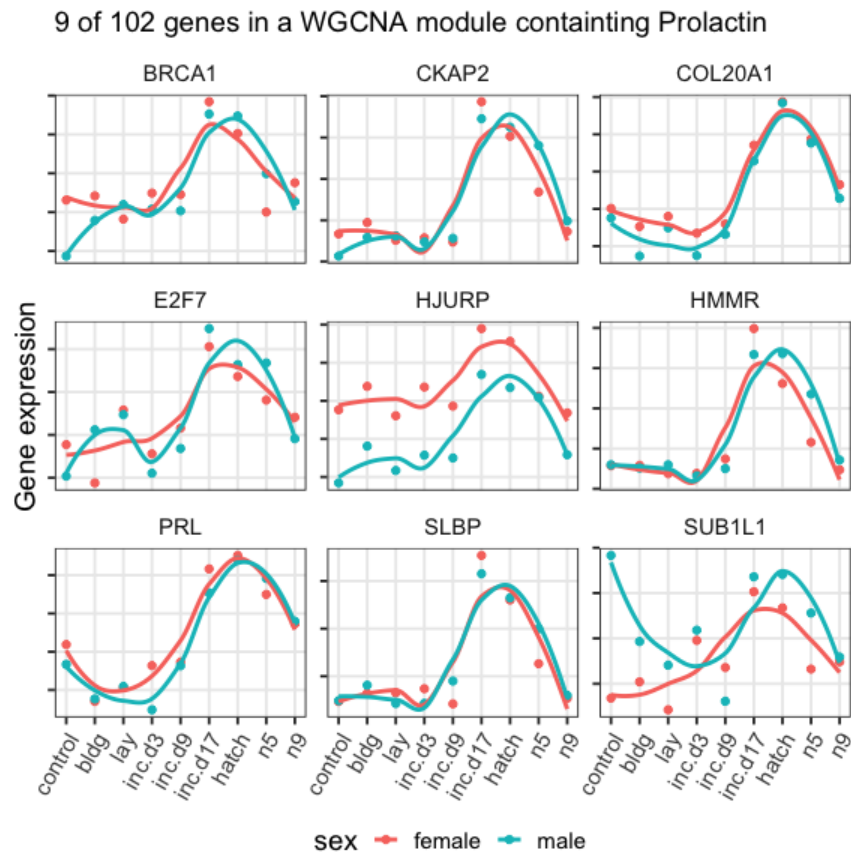


Fig 4: Hundreds of genes show similar pattern in expression as circulating prolactin



Acknowledgments

This project is a synergistic collaboration between the PI, Rebecca Calisi-Rodríguez (expertise in avian behavior, parental care and neurobiology), co-PI, Matthew MacManes (expertise in next-generation sequencing, transcriptome assembly, and gene expression analyses), and Collaborator, Rae Silver (expertise in neurobiology, dove behavior, and decades of successful breeding and maintenance of dove colonies at Barnard College).

References

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