Comparative dynamics of female germ cell populations : insight from imaging and multiscale modeling

Frédérique Clément, Romain Yvinec

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Journées INRAE-INRIA, 04-05 Juillet 2023

Outline

Collaborative background

Biological background

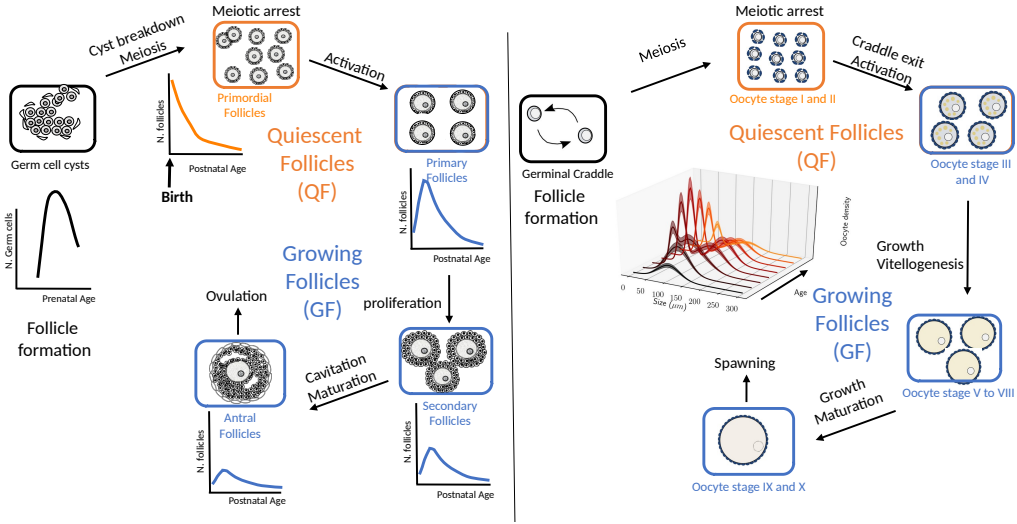
Available and future data

Modeling questions and approaches

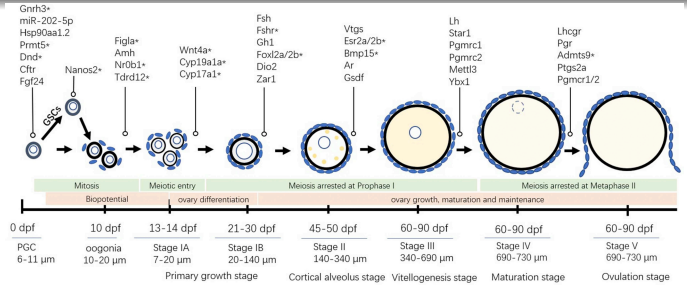
Collaborative background

- EPC CNRS-INRAE-INRIA MUSCA
  MUltiSCAle population dynamics for physiological systems
  CRI Saclay – MaiAGE – PRC
- Projet GinFiz ANSES 2020
  Gonadal aromatase inhibition and other toxicity pathways leading to Fecundity Inhibition in Zebrafish: from initiating events to population impacts
  collaboration INERIS (Rémy Beaudouin) + Laboratoire de Physiologie et Génomique des Poissons (LPGP, Violette Thermes)
- Projet IMMO Digit-Bio INRAE 2021
  Imagerie et modélisation multi-échelles pour la compréhension de la dynamique ovarienne chez le poisson
  collaboration LPGP
- AAPG ANR CES 45 OVOPAUSE 2022
  Dynamics and regulation of female germ cell populations: understanding aging through population dynamics models
  collaboration LPGP + INSERM

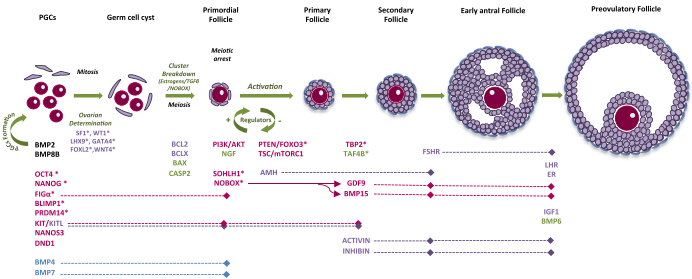
Comparative vertebrate oogenesis (1)



Comparative vertebrate oogenesis (2)



Li & Ge Mol. Cell. Endocrinol. 2020



Sánchez & Smith Acta Bioch. Biophys. 2012

Main questions and related outcomes

Population scale

- Kinetics of oocyte pool exhaustion / intensity of oocyte pool renewal
- Shaping of the oocyte (size/maturation) distribution
- Contribution of direct and indirect interactions within the oocyte population

  Management of oocyte resources / Driving of ovarian cyclicity

Oocyte/follicle scale

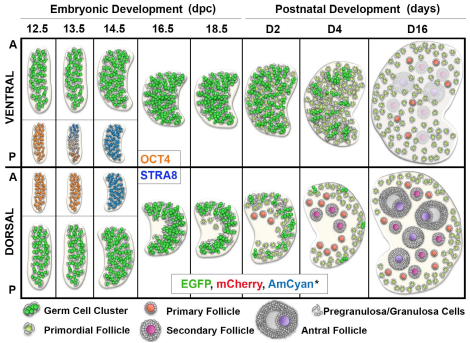
- Coupled dynamics between germ cells and somatic cells
- Mechanisms underlying the proper sequence of morphogenetic events

Preserving the ovarian resources

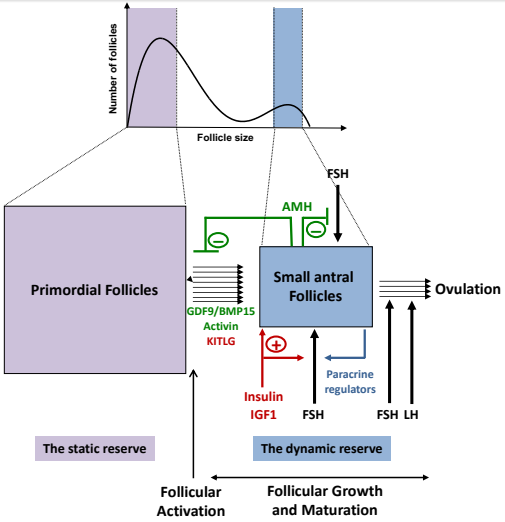
- Ovarian aging
- Reproductive fitness

Knowledge driven modeling approaches (Mammals)

Embedding cell biology/developmental biology/endocrine information



Cordeiro et al. Biol. Reprod. 2015



Monniaux et al. Front. Endocr. Dis. 2019

Knowledge driven modeling approaches (Mammals)

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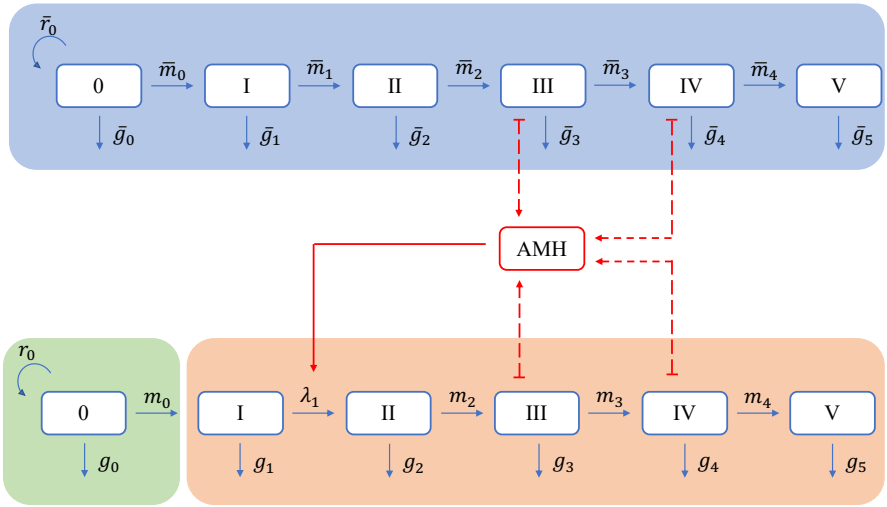
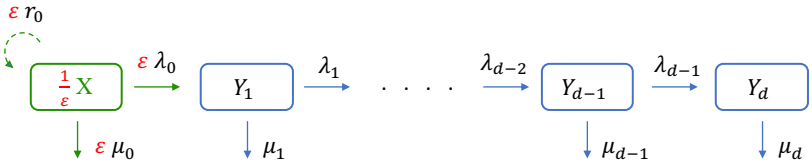


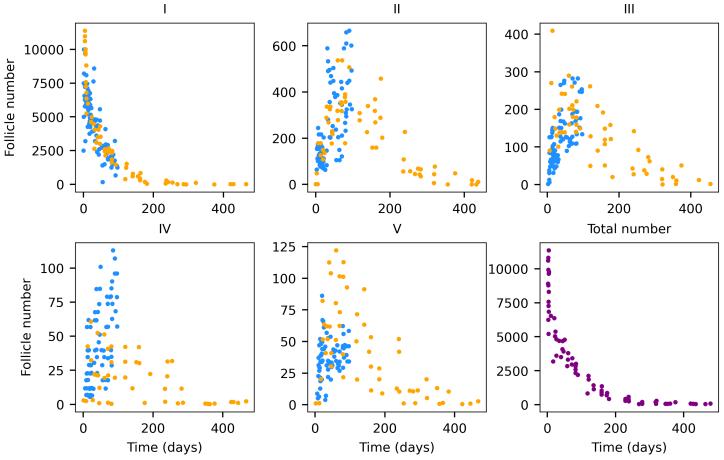
Stochastic compartmental population dynamics

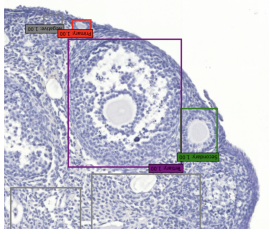
Multiple timescales and order of magnitudes $\Rightarrow$ Model reduction



	Transition	Rate
Birth (reserve)	$\left(X^\varepsilon, Y^\varepsilon\right) \to \left(X^\varepsilon + \varepsilon, Y^\varepsilon\right)$	$\frac{r_0(Y^\varepsilon)}{\varepsilon} X^\varepsilon$
Maturation (reserve)	$\left(X^\varepsilon, Y^\varepsilon\right) \to \left(X^\varepsilon - \varepsilon, Y^\varepsilon + e_1\right)$	$\frac{\lambda_0(Y^\varepsilon)}{\varepsilon} X^\varepsilon$
Death (reserve)	$\left(X^\varepsilon, Y^\varepsilon\right) \to \left(X^\varepsilon - \varepsilon, Y^\varepsilon\right)$	$\frac{\mu_0(Y^\varepsilon)}{\varepsilon} X^\varepsilon$
Maturation, $i \in [\![1, d-1]\!]$	$\left(X^\varepsilon, Y^\varepsilon\right) \to \left(X^\varepsilon, Y^\varepsilon - e_i + e_{i+1}\right)$	$\frac{\lambda_i(Y^\varepsilon)}{\varepsilon} Y_i^\varepsilon$
Death, $i \in [\![1, d]\!]$	$\left(X^\varepsilon, Y^\varepsilon\right) \to \left(X^\varepsilon, Y^\varepsilon - e_i\right)$	$\frac{\mu_i(Y^\varepsilon)}{\varepsilon} Y_i^\varepsilon$

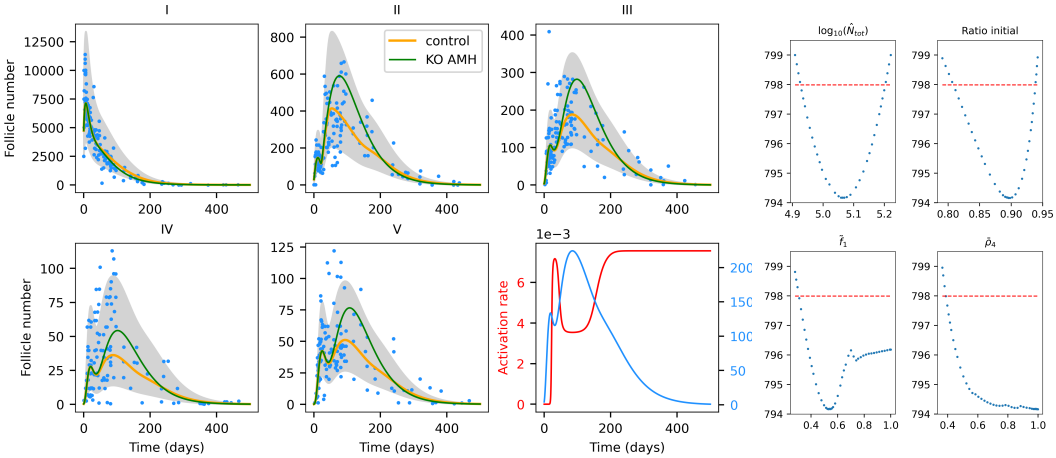
Data-driven parameter estimation : low-throughput data





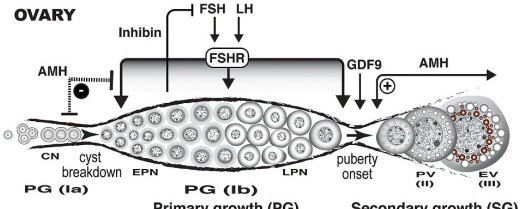
Data-driven parameter estimation : low-throughput data

Model selection, parameter identifiability, perturbation prediction

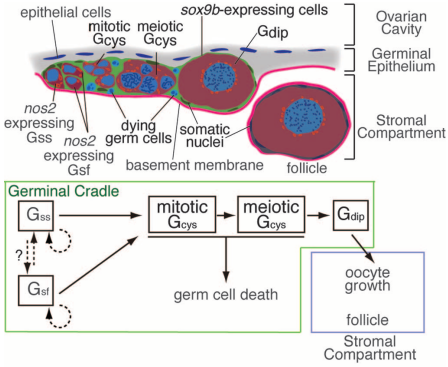


Knowledge driven modeling approaches (Fish)

Embedding cell biology/developmental biology/endocrine information



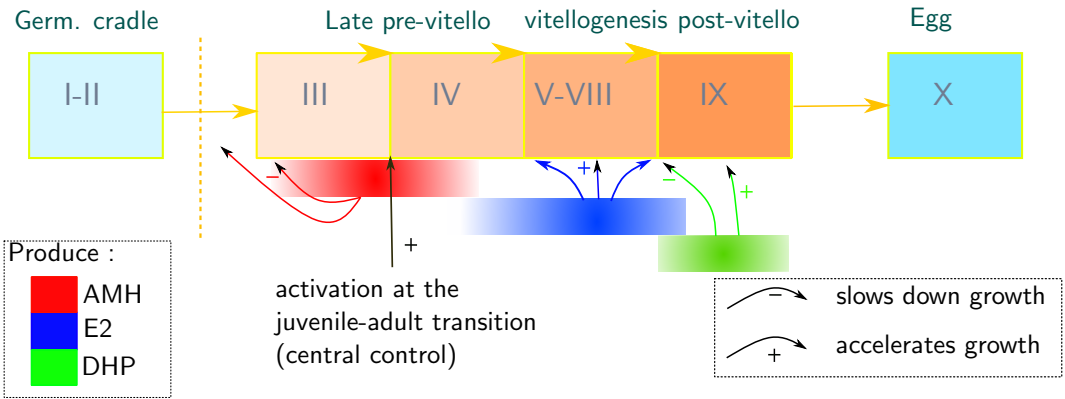
Zhang et al. Mol. Cell. Endocrinol. 2020



Nakamura et al. Science 2010

Knowledge driven modeling approaches (Fish)

Embedding cell biology/developmental biology/endocrine information

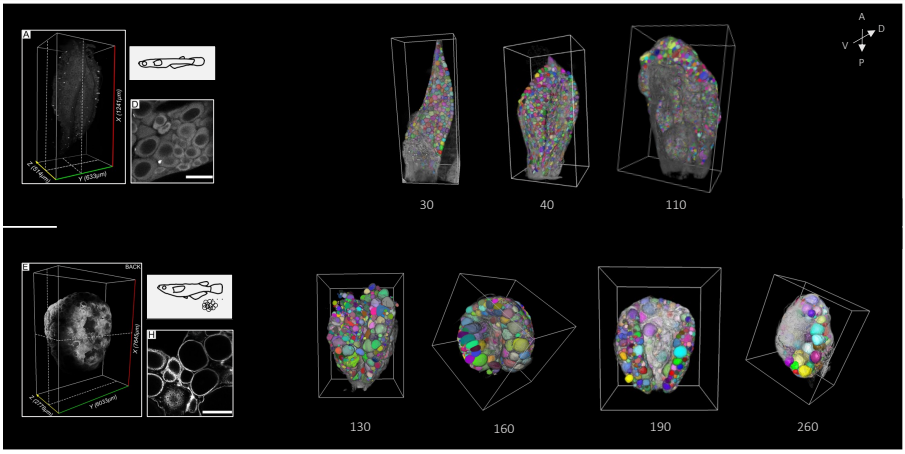


Parameters	Interpretation	**Output**	Interpretation
λ_0	Cradle exit rate	ρ_0	number of cells in the cradle
r_0	Cradle renewal rate	ρ	size density from stage III to IX
λ	growth speed from stage III to IX	ρ_1	number of stage X oocytes
W_i	"quantity" of hormone i secreted		

$$
\begin{cases}
\frac{d}{dt}\rho_0(t) = r_0(\rho_0)\rho_0(t) - \lambda_0(W_{AMH}(t))\rho_0(t), \ t > 0 \\
\lim\limits_{x\to 0}(\lambda\rho) = \lambda_0\rho_0(t), \quad \text{sur } [0,+\infty) \\
\partial_t\rho + \partial_x\left(\lambda(x, W_{AMH}, W_{E2}, W_{DHP})\rho\right) = 0, \quad x \in [0,1], \ t > 0 \\
\frac{d}{dt}\rho_1(t) = \lim\limits_{x\to 1}(\lambda\rho) - \text{spawn}(t), \ t > 0 \\
W_i(t) = \int_0^1 \omega_i(x)\rho(t,x)\mathrm{d}x, \quad i \in \{AMH, E2, DHP\}
\end{cases}
$$

Data-driven parameter estimation : DL-based data extraction

Work of Violette Thermes and collaborators

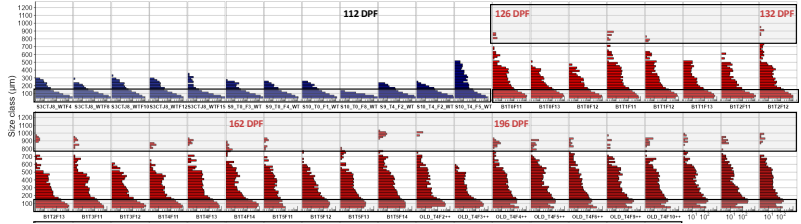


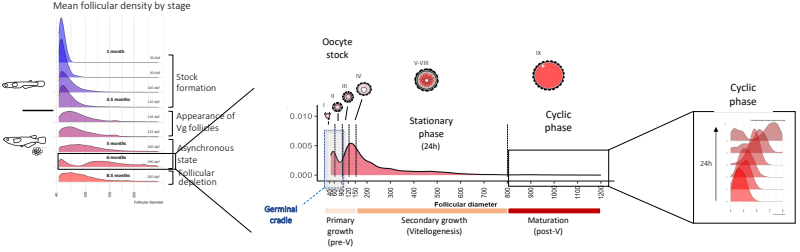
Inputs : 3D ovarian imaging / Automatic follicle segmentation and classification

Outputs : age/space-varying distribution in size/class of the total population of ovarian follicles

Data-driven parameter estimation : DL-based data extraction

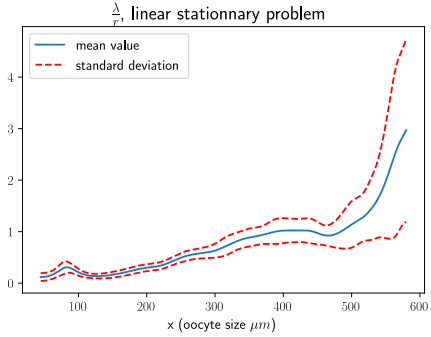
Work of Violette Thermes and collaborators

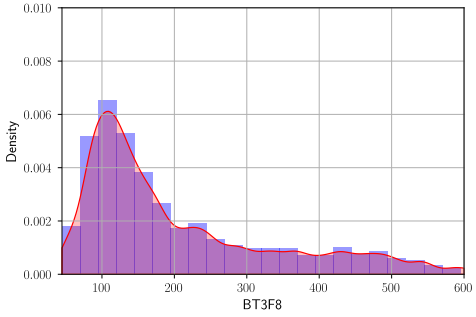




$$\begin{cases} \bar{\rho}(0) = r \\ \partial_x \left(\lambda(x)\bar{\rho}\right) = 0, \quad x \in [0,1] \end{cases}$$

Hormonal interactions cannot be deduced from purely stationary data, yet we can infer the size-dependent oocyte growth speed.





Ongoing/ future directions

Stochastic and deterministic models of structured populations with nonlinear and nonlocal terms

- Wellposedness / stationary solutions
- Inverse problems
- Structuring variable(s)
  Coupling with cell dynamics models on the single-follicle level
  Spatial distribution
- Physics-based modeling (morphogenesis)