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Time scale separation in life-long ovarian follicles population dynamics model

Romain Yvinec

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Submitted on 1 Sep 2023

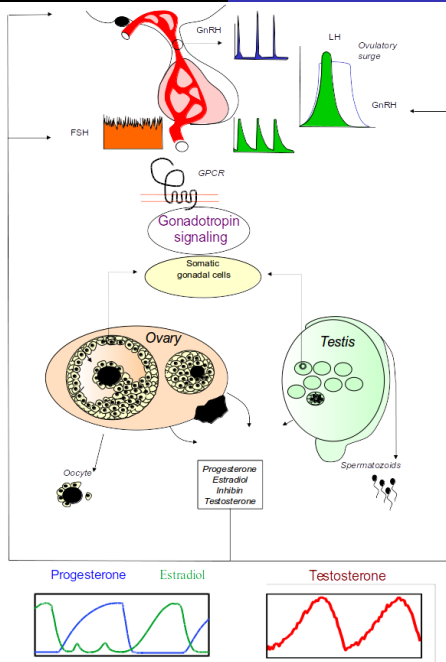
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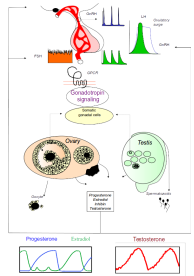
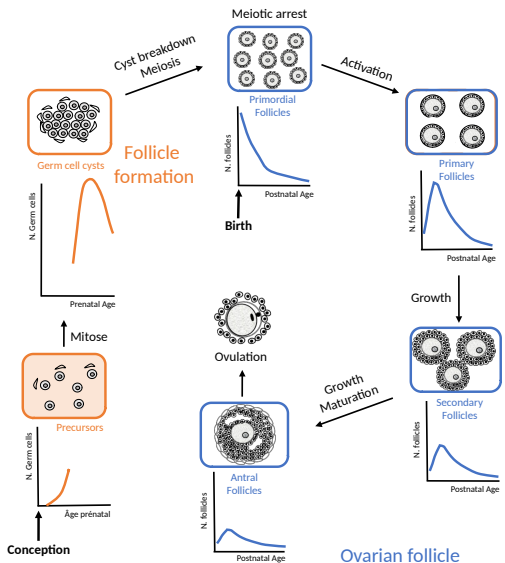
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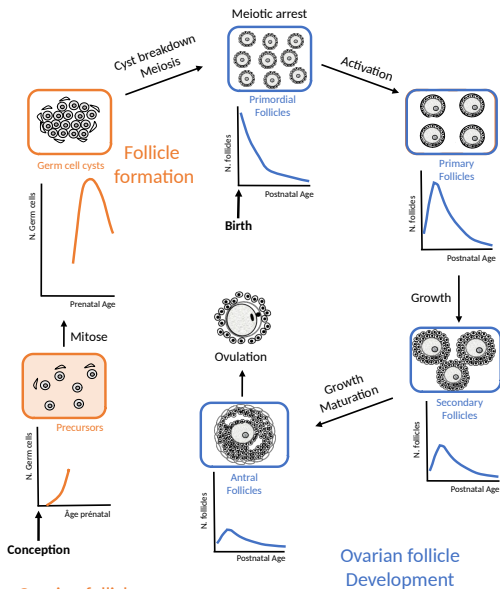
BIOS Team, Physiologie de la Reproduction et des Comportements, INRAE CNRS,
Université de Tours, (Tours, France)
MUSCA Team, INRIA-INRAE-CNRS (Saclay, France)





Ovarian follicle Formation

Ovarian follicle Development



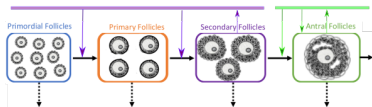
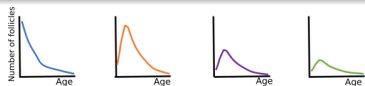
Ovarian follicle Formation

Ovarian follicle Development

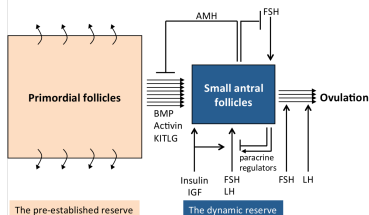
- Pool of Quiescent follicles **static reserve** (perinatal in most mammals)
Slow activation
- Basal growth **Dynamic reserve** (starting at birth) Spanning over several ovarian cycles
- Terminal growth
After puberty : **ovulation** within an ovarian cycle

Population dynamics in female gametogenesis

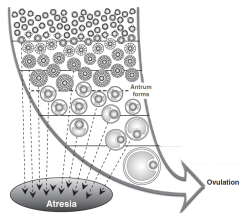
- Asynchronous growth
- Interactions between subpopulations



Ovarian reserves of follicles and their regulations



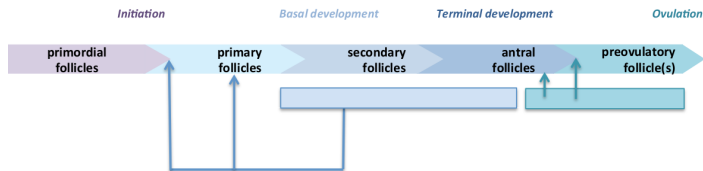
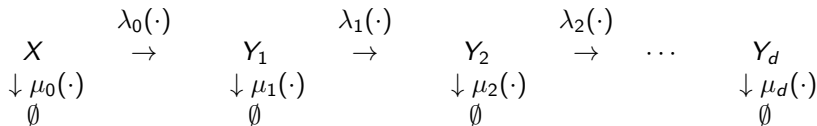
Monniaux, *Theriogenology* 2016



Scaramuzzi et al., *Reprod.Fert. Dev.* 2011

Lifespan follicle population model

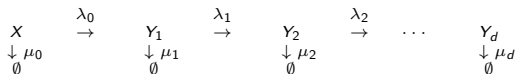
- Structured Population in compartments
- Non linear interaction between follicles *via* λ' 's and μ' 's.



Bonnet et al. *Multiscale population dynamics in reproductive biology : singular perturbation reduction in deterministic and stochastic models*, ESAIM : PROCEEDINGS AND SURVEYS, 2020.

Lifespan follicle population model

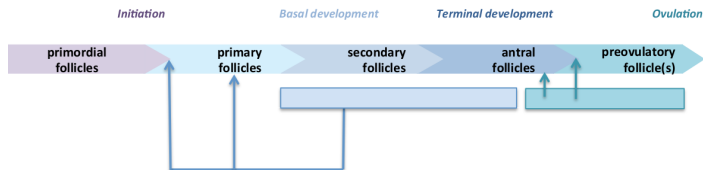
- Structured Population in compartments
- Non linear interaction between follicles *via* λ' 's and μ' 's.



Typical choice

$$\lambda_i(Y) = m_i + \frac{f_i}{1 + K_{1,i} \sum_{j=1}^d \omega_{1,j} Y_j},$$

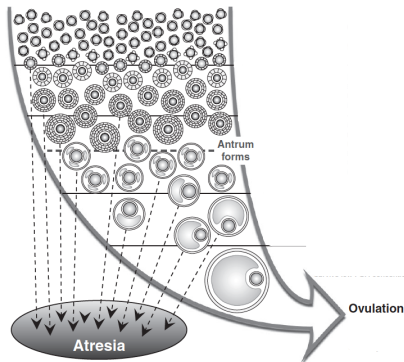
$$\mu_i(Y) = g_i \left(1 + K_{2,i} \sum_{j=0}^d \omega_{2,j} Y_j \right)$$



Bonnet et al. *Multiscale population dynamics in reproductive biology : singular perturbation reduction in deterministic and stochastic models*, ESAIM : PROCEEDINGS AND SURVEYS, 2020.

Order of magnitude (in Women)

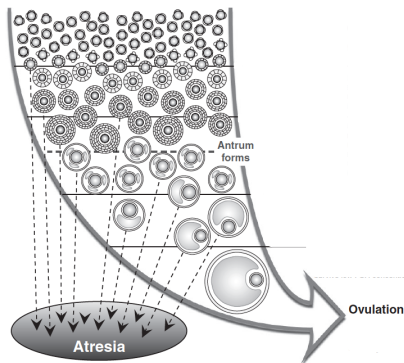
- Quiescent follicles
 - peri-natal $\approx 5 \cdot 10^6$
 - At birth $\approx 1 \cdot 10^6$
 - At puberty $10^4 - 10^6$
 - At menopause $< 10^3$
 - Activation rate "A few per days"



Scaramuzzi et al., *Reprod.Fert. Dev.* 2011

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 - Activation rate "A few per days"
- Growing follicles
 - Maturation time 120 – 180j
 - Basal follicles $10^3 - 10^4$
 - Terminal follicles 10^2
 - Pre-Ovulatory follicles a few
 - Atresia Most of them



Scaramuzzi et al., *Reprod.Fert. Dev.* 2011

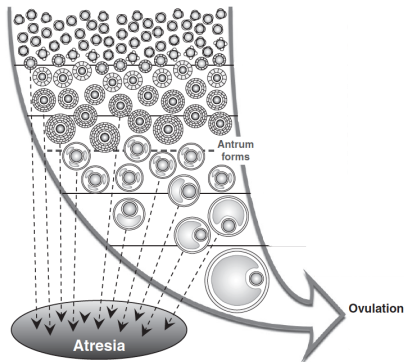
Order of magnitude (in Women)

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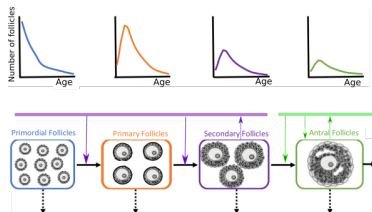
peri-natal	$\approx 5 \cdot 10^6$
At birth	$\approx 1 \cdot 10^6$
At puberty	$10^4 - 10^6$
At menopause	$< 10^3$
Activation rate	"A few per days"
- Growing follicles

Maturation time	120 – 180j
Basal follicles	$10^3 - 10^4$
Terminal follicles	10^2
Pre-Ovulatory follicles	a few
Atresia	Most of them

> **Only 400 follicles will ever reach the pre-ovulatory stage**

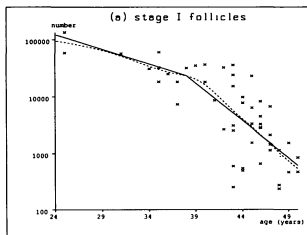


Scaramuzzi et al., *Reprod.Fert. Dev.* 2011

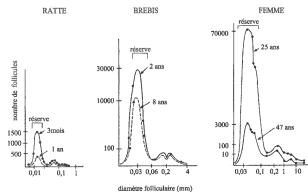


⇒ Irreversible (slow) decay of an initial pool of quiescent follicle

⇒ "Stable" repartition of growing follicle

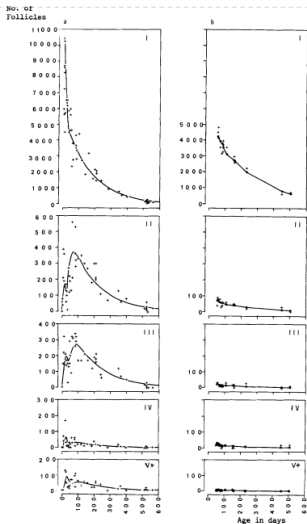


Faddy and Gosden 1995



Thibault and Levasseur, 2001

Previous compartmental model (mice data)

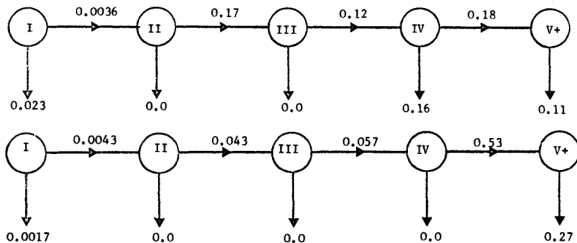


Faddy et al., J. Exp. Zool. 1976

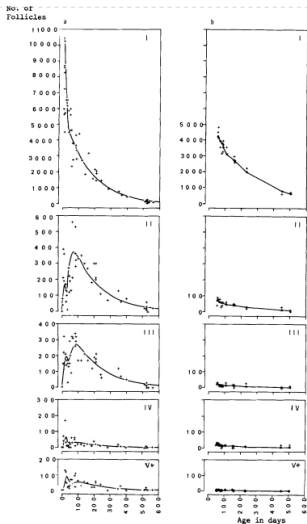
An Analytical Model for Ovarian Follicle Dynamics

M. J. FADDY,¹ ESTHER C. JONES² AND R. G. EDWARDS³
¹ Department of Mathematical Statistics, University of Birmingham, Birmingham B15 2TT, U.K., ² Department of Anatomy, University of Birmingham, Birmingham B15 2TJ, U.K., and ³ Physiological Laboratory, University of Cambridge, Cambridge CB2 3EG, U. K.

- "Migration-death" model
- Linear Model



Previous compartmental model (mice data)

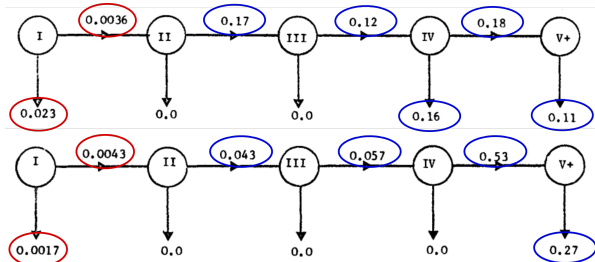


Faddy et al., J. Exp. Zool. 1976

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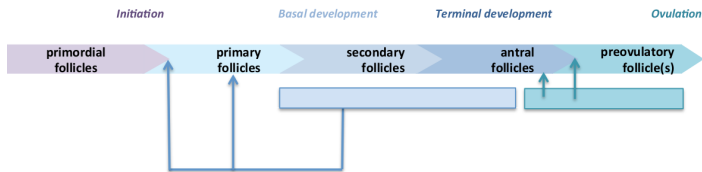
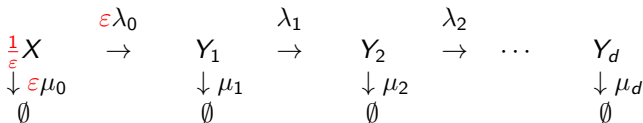
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Lifespan follicle population model

- Structured Population in compartments
- Non linear interaction between follicles *via* λ 's and μ 's.
- Two time and abundance scales ($\varepsilon \ll 1$)

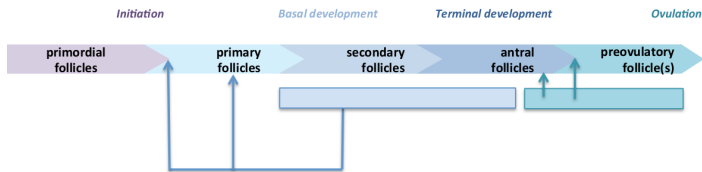
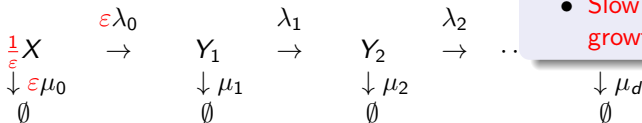


Lifespan follicle population model

- Structured Population in compartments
- Non linear interaction between follicles *via* λ 's and μ 's.
- Two time and abundance scales ($\varepsilon \ll 1$)

Time scale separation

- Quiescent Pool \gg Growing Follicles
- Slow Activation \ll Fast growth



Singular Perturbation Theory

$$\begin{array}{ccccccc}
 & \varepsilon \lambda_0 & & \lambda_1 & & \lambda_2 & & \dots & & Y_d \\
 \frac{1}{\varepsilon} X & \rightarrow & Y_1 & \rightarrow & Y_2 & \rightarrow & \dots & & & Y_d \\
 \downarrow \varepsilon \mu_0 & & \downarrow \mu_1 & & \downarrow \mu_2 & & & & & \downarrow \mu_d \\
 \emptyset & & \emptyset & & \emptyset & & & & & \emptyset
 \end{array}$$

In the limit $\varepsilon \rightarrow 0$ and in the time scale t/ε :

We expect X and $Y = (Y_1, \dots, Y_d)$ to converge to a differential-algebraic equation :

$$\begin{cases} \frac{dx}{dt}(t) = F(x(t), y(t)), & x(0) = x^{\text{in}}, \\ 0 = G(x(t), y(t)), & t > 0 \end{cases}$$

$$\begin{array}{ccccccc}
 & \varepsilon \lambda_0 & & \lambda_1 & & \lambda_2 & & \dots & & Y_d \\
 \frac{1}{\varepsilon} X & \rightarrow & Y_1 & \rightarrow & Y_2 & \rightarrow & \dots & & & Y_d \\
 \downarrow \varepsilon \mu_0 & & \downarrow \mu_1 & & \downarrow \mu_2 & & & & & \downarrow \mu_d \\
 \emptyset & & \emptyset & & \emptyset & & & & & \emptyset
 \end{array}$$

Theorem (G. Ballif, F. Clément, R.Y. SIAP 2022)

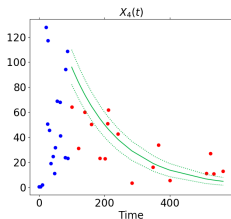
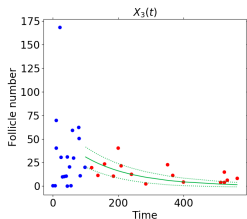
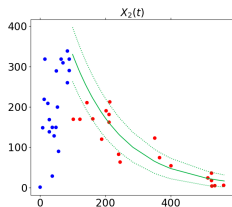
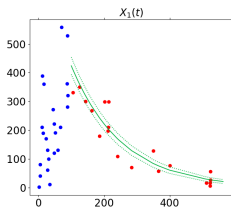
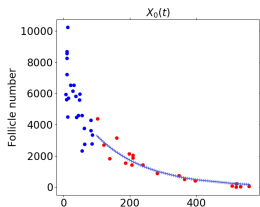
(...) $(X^\varepsilon, Y^\varepsilon)$ converges in $\mathcal{D}_{\mathbb{R}}[0, \infty[\times \mathcal{L}_m(\mathbb{N}^d)$ to the unique solution of

$$\begin{cases} \frac{dx}{dt}(t) & = \Lambda_0(x(t))x(t), & x(0) = x^{\text{in}}, \\ \Lambda_0(x(t)) & = - \sum_{y \in \mathbb{N}^d} (\lambda_0(y) + \mu_0(y)) \pi_{x(t)}(y), \end{cases}$$

$$\sum_{y \in \mathbb{N}^d} L_x \psi(y) \pi_x(y) = 0, \quad \forall \psi \text{ bounded on } \mathbb{N}^d,$$

$$\begin{aligned}
 L_x \psi(y) = \lambda_0(y) x & \left[\psi(y + e_1) - \psi(y) \right] + \sum_{i=1}^{d-1} \lambda_i(y) y_i \left[\psi(y + e_{i+1} - e_i) - \psi(y) \right] \\
 & + \sum_{i=1}^d \mu_i(y) y_i \left[\psi(y - e_i) - \psi(y) \right].
 \end{aligned}$$

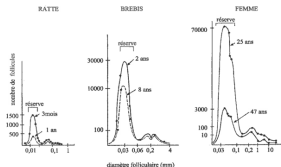
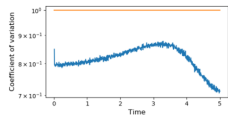
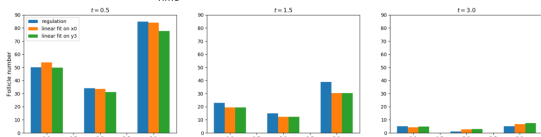
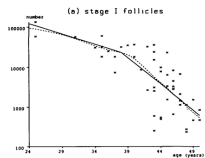
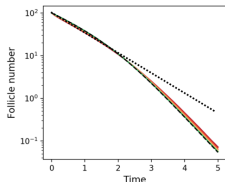
Does it works in practice ?



$$\begin{cases} x_0(t) &= X_0^{in} \exp\left(-(\lambda_0 + \mu_0)t\right) \\ \pi_{x_0(t)}(y) &= \prod_{i=1}^d \frac{(x_0(t)\alpha_i)^{y_i}}{y_i!} e^{-x_0(t)\alpha_i} \\ \alpha_i &= \prod_{j=0}^{i-1} \frac{\lambda_j}{\lambda_{j+1} + \mu_{j+1}} \end{cases}$$

- Timescale separation is coherent with published data on follicle counts in mice at the lifespan time scale, *away from a transient period*.

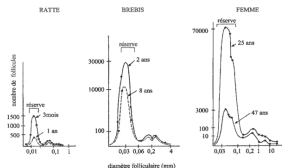
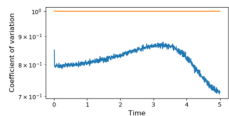
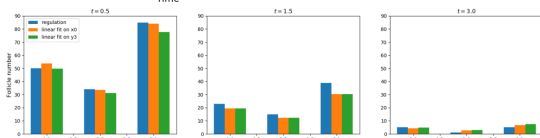
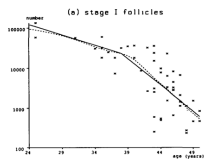
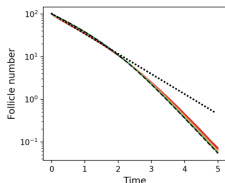
What is it useful for?



- ✓ acceleration of reserve decay with age :

$$\frac{dx}{dt} = -\left(a + \frac{b}{1+cx}\right)x$$
 -> Mechanistic explanation of previously published statistical regression model (Coxworth and Hawkes 2010)

What is it useful for?



- ✓ acceleration of reserve decay with age :

$$\frac{dx}{dt} = -\left(a + \frac{b}{1+cx}\right)x$$
 -> Mechanistic explanation of previously published statistical regression model (Coxworth and Hawkes 2010)
- ✓ "stable" evolution of growing follicles Y
 -> Antral Follicle Count for fertility test and primary ovarian insufficiency detection

Going further

- What about fluctuations ?
- Can we infer the regulation mechanism that control follicle activation ?
- Can we refine the model to model the transient phase (reserve establishment / early post-natal dynamics) ?

Fluctuations

At leading order, fluctuations are Gaussian of order $\sqrt{\varepsilon}$: $U^\varepsilon = \frac{X^\varepsilon - x}{\sqrt{\varepsilon}}$

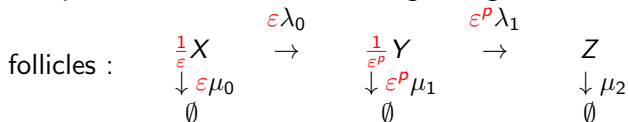
CLT

... U^ε converges towards U that satisfies

$$U(t) = U^{in} + \int_0^t \left[\lambda'_0(x(s))x(s) + \lambda(x(s)) \right] U(s) ds + \int_0^t \sqrt{\langle G(x(s), \cdot), \pi_{x(s)} \rangle} W_s$$

Fluctuations with more than two timescales

X =quiescent follicles; Y =small growing follicles; Z large terminal

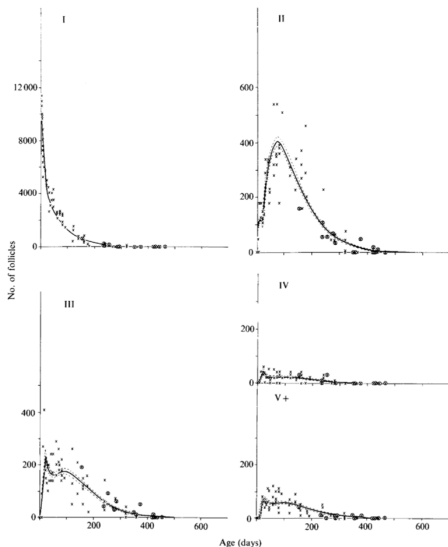


For $p > 1/2$, fluctuations on X are still on order $\sqrt{\varepsilon}$, with Y and Z contributing *equally*.

For $p \leq 1/2$: undergoing work !

"Time" course

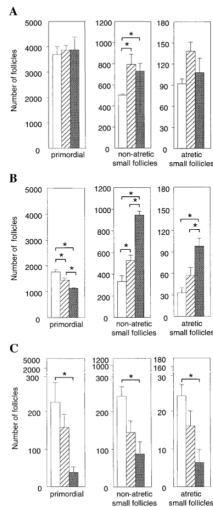
- Follicle count in mice from birth until 500 days.
- Reserve + 4 compartments (Faddy's classification)
- (Recovery of points by hand)



Faddy, Gosden and Edwards, J. Endo., 1983
 Faddy, Telfer and Gosden, Cell Proliferation, 1987

Perturbation data : KO AMH

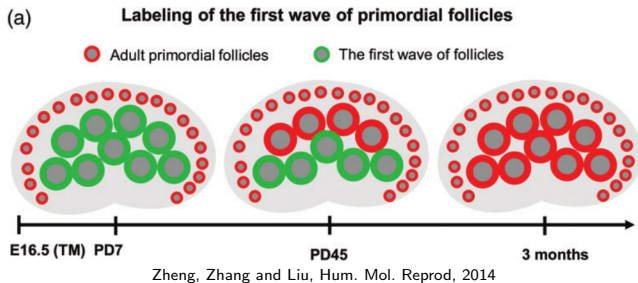
- AMH Inhibition in vivo on mice
- 3 genotypes : control group (+/+), heterozygous mice KO AMH (+/-) homozygous mice KO AMH (-/-).
- Follicle counts at 3 ages :
 - 25 days (A)
 - 120 days (B)
 - 390 days (C)



Durlinger and al, Endocrinology, 1999

Ovarian Reserve build-up

- Two distinct population of follicles are present initially
- Labelling of each population

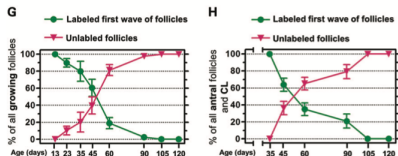
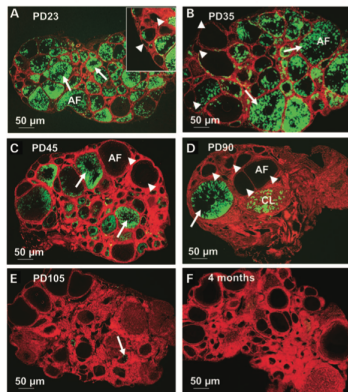


Ovarian Reserve build-up

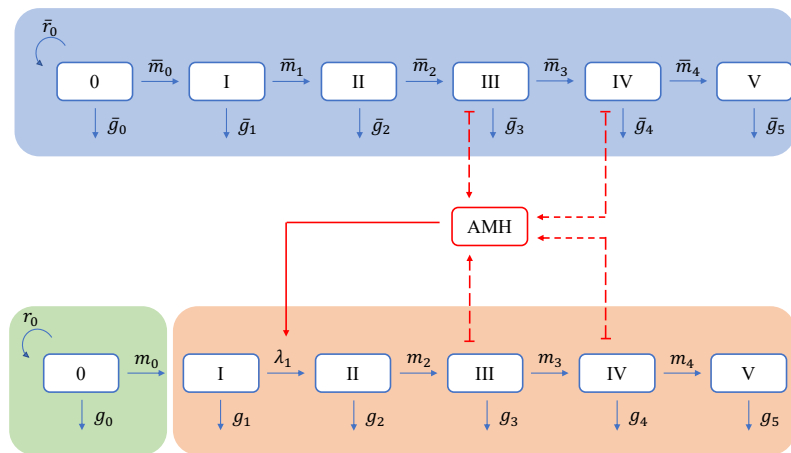
- Tracing follicles of the first wave of activated follicles.
- Proportion of first wave activated follicles among growing follicles.

$$p(t) = \frac{\sum_{i=1}^4 X_i^1(t)}{\sum_{i=1}^4 X_i^{tot}(t)}$$

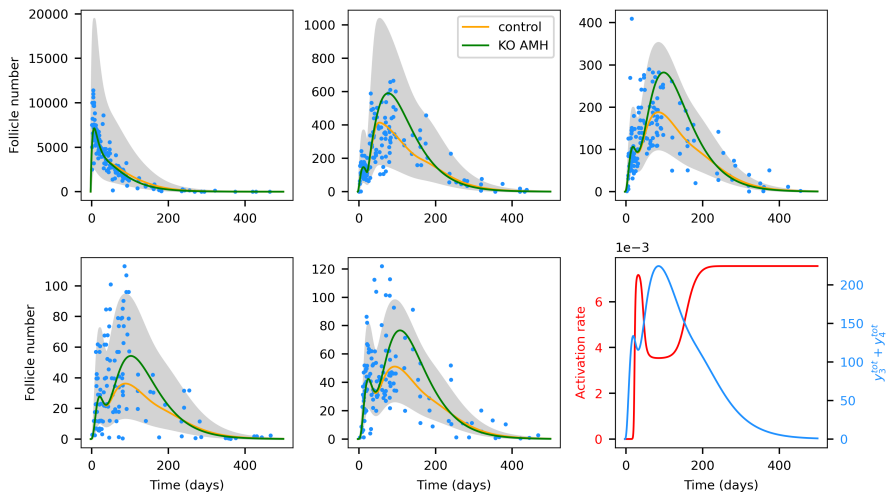
Tamoxifen was given at E16.5 and ovaries were analyzed at various ages



ODE model

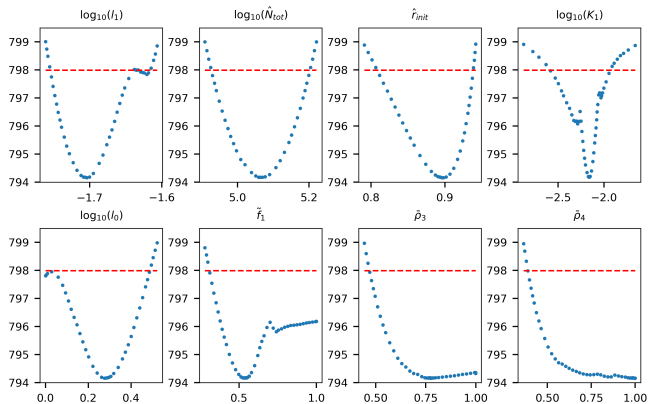


Data fitting



Identifiability

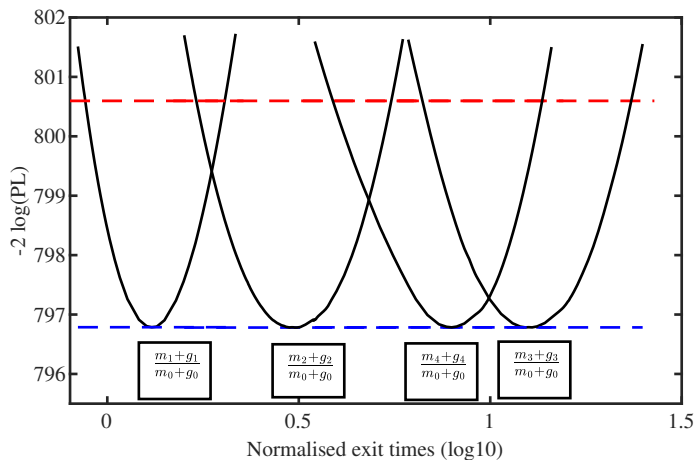
Theoretical (*Structural Identifiability Julia package*) and practical identifiability (*Data2Dynamics*)



Matlab p. 14

- 20 of 31 parameters are practically identifiable.
- Additional Data on germ cell dynamics would greatly improve identifiability

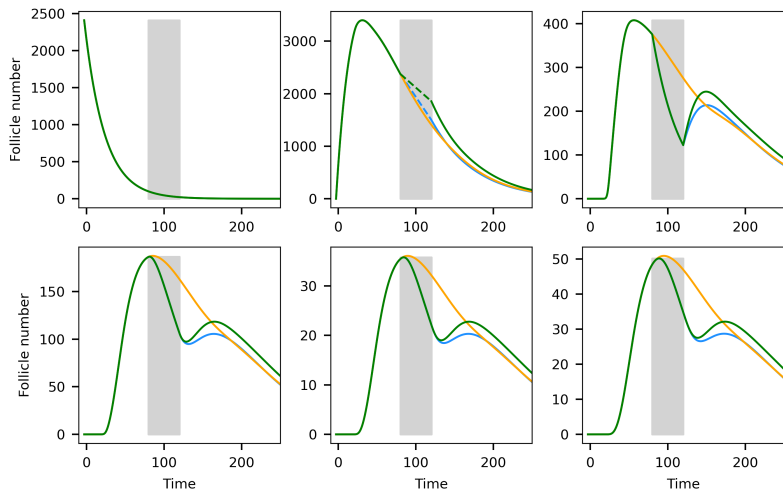
Timescale difference between growing / quiescent follicle



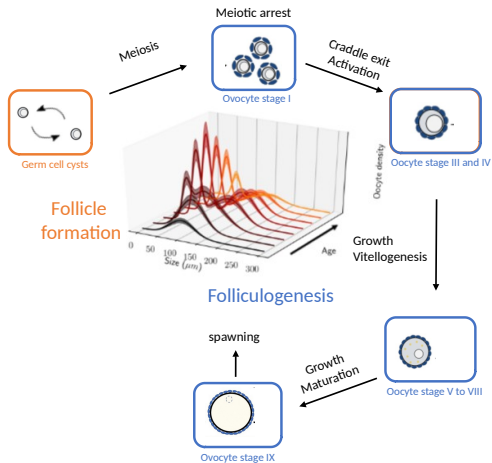
Parameter values interpretation

- Mean activation time is around 200 days, while growing time is around 50 days.
- The first-wave follicles is approx. 5 times faster than the second wave for compartments 0,1,2
- Follicles atresia is negligible in compartment 2,3,4

Prediction of AMH administration

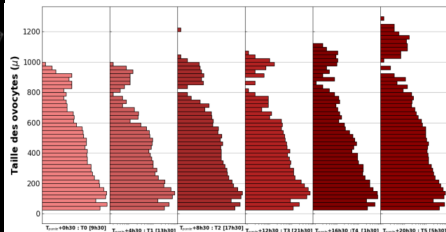
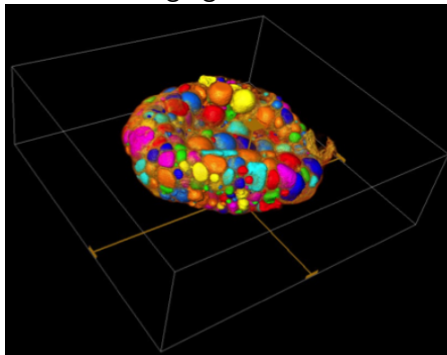


Comparison with fish oogenesis



Continuous structuring variable

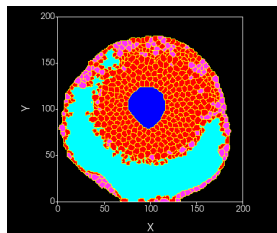
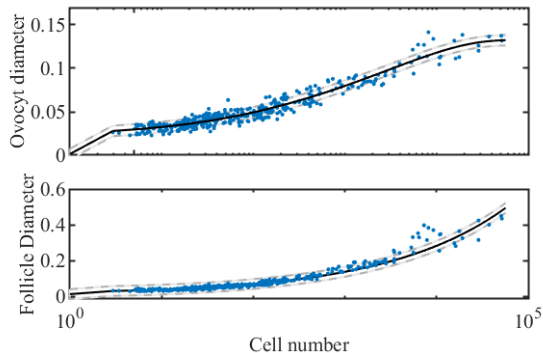
3D imaging data : whole Follicle count and size measurement



-> The bulk size distribution is ovarian-cycle independent (slow variable).

$$\begin{cases} \frac{d\rho_0(t)}{dt} &= -(\lambda_0(\rho(t, \cdot)) + \mu_0(\rho(t, \cdot)))\rho_0(t), \\ \varepsilon \partial_t \rho(t, x) &= -\partial_x (\lambda(\rho(t, \cdot), x)\rho(t, x)) - \mu(\rho(t, \cdot), x)\rho(t, x), \quad \text{for} \\ \lim_{x \rightarrow 0} \lambda(\rho(t, \cdot), x)\rho(t, x) &= \lambda_0(\rho(t, \cdot))\rho_0(t), \end{cases}$$

More than one structuring variable



Conclusion and perspective

- ✓ Lifespan ovarian follicle population dynamics model
- ✓ Separation of time scale explains slow decay of the reserve and quasi stable growing follicle repartition
- ✓ Follicle count data and perturbation experiments may reveal feedback mechanisms
 - Extension to three timescale (reserve, basal growth and terminal growth)
 - Extension to (several) continuous structuring variable
 - Comparative physiology approach

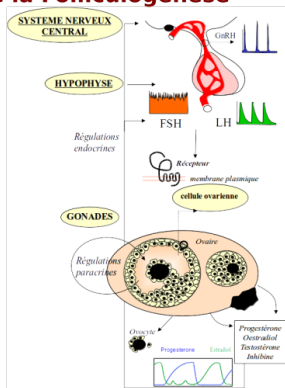
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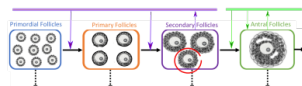
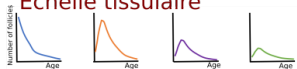
Open Post-position available in 2023 !

Thanks for your attention !

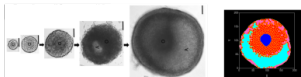
Modélisation Multi-échelle de la Folliculogénèse



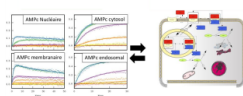
Echelle tissulaire



Echelle cellulaire



Echelle intra-cellulaire



Inria INRAE

- * INRIA Saclay : Frédérique Clément, Guillaume Ballif, Frédérique Robin
- * INRAE PRC : Team BIOS, BINGO (Danielle Monniaux, Véronique Cadoret, Rozenn Dalbies-Tran)
- * INRAE LPGP (Violette Thermes)
- * INSERM - Paris Cité (Céline Guigon)
- * Lea Popovic (Concordia University, Montreal)
- * CEMRACS 2018 (Céline Bonnet, Keltoum Chahour)