



# Reconstructing the origins of animal multicellularity using experimental evolution

## Fact Sheet

### Project Information

#### MULTICELLEXPVO

Grant agreement ID: 746044

[Project website](#)

#### DOI

[10.3030/746044](https://doi.org/10.3030/746044)

Project terminated on 28 February 2020

#### EC signature date

17 March 2017

#### Start date

1 September 2018

#### End date

31 August 2020

#### Funded under

EXCELLENT SCIENCE - Marie Skłodowska-Curie Actions

#### Total cost

€ 158 121,60

#### EU contribution

€ 158 121,60

#### Coordinated by

AGENCIA ESTATAL CONSEJO  
SUPERIOR DE  
INVESTIGACIONES CIENTIFICAS  
 Spain

## Objective

All living animals are descended from a single-celled ancestor. Understanding how this ancestor became the first multicellular animal remains a major challenge in the field of evolutionary biology. Phylogenomic analyses have shown that animals are closely related to three unicellular lineages: choanoflagellates, filastereans and ichthyosporeans, altogether forming the Holozoa clade. Genetic and phenotypic studies have shown that the filasterean *Capsaspora owczarzaki* can under specific growth conditions form transient multicellular aggregates. However, why is this

multicellularity only transient? What are the genetic and phenotypic requirements for its emergence and stabilization? And what is the role of the actin cytoskeleton in this transition? Indeed the actin cytoskeleton is known for its pivotal role for cell coordination and morphology, which must play a role in evolution of multicellularity. To address these questions, we will use the *C. owczarzaki* as a model organism. We will combine cell biology, genomics and experimental evolution to unravel multicellularity emergence and stabilization. Specifically, we will aim to obtain evolved mutants showing excessive and more stable multicellular behaviour of *C. owczarzaki* using long-term experimental evolution. Such evolved strains would unravel how multicellularity emerged and stabilized. In addition, using random mutagenesis screen, we aim to identify mutants unable to form multicellular aggregates. Such mutants would reveal the minimum genetic requirements for such a transition. Finally, we will take advantage of recently developed genetic tools in *C. owczarzaki* to study the actin cytoskeleton during the cell cycle. Our results could reveal how the first multicellular ancestor of animals appeared from a genetic and cellular perspective, and, how cell fate specification was established during evolution. Results generated on this fellowship will be relevant to evolutionary, cell and developmental biologists.

## Fields of science (EuroSciVoc)

[natural sciences](#) > [biological sciences](#) > [genetics](#)

[natural sciences](#) > [biological sciences](#) > [evolutionary biology](#)

[natural sciences](#) > [biological sciences](#) > [cell biology](#)



## Keywords

[Multicellularity emergence](#)

[Metazoa](#)

[Actin organization](#)

[Experimental evolution](#)

## Programme(s)

[H2020-EU.1.3. - EXCELLENT SCIENCE - Marie Skłodowska-Curie Actions](#)

MAIN PROGRAMME

[H2020-EU.1.3.2. - Nurturing excellence by means of cross-border and cross-sector mobility](#)

## Topic(s)

## Call for proposal

[H2020-MSCA-IF-2016](#)

[See other projects for this call](#)

## Funding Scheme

[MSCA-IF - Marie Skłodowska-Curie Individual Fellowships \(IF\)](#)

## Coordinator



### AGENCIA ESTATAL CONSEJO SUPERIOR DE INVESTIGACIONES CIENTIFICAS

Net EU contribution

**€ 158 121,60**

Total cost

**€ 158 121,60**

Address

**CALLE SERRANO 117**

**28006 Madrid**

 **Spain** 

Region

**Comunidad de Madrid > Comunidad de Madrid > Madrid**

Activity type

**Research Organisations**

Links

[Contact the organisation](#)  [Website](#) 

[Participation in EU R&I programmes](#) 

[HORIZON collaboration network](#) 

**Last update:** 23 July 2023

**Permalink:** <https://cordis.europa.eu/project/id/746044>

