ZooSize_SOP_v1.0 Last edited: 2021-11-19

Standard operating protocol for measuring crustacean zooplankton body length

This SOP has been created for the **ZooSize¹ project**. References and methods from colleagues² have been combined by the ZooSize champions³ to produce a general guideline for subsampling and measurements of individual zooplankton species body lengths.

Collaborators who have not yet analysed their sample can choose to follow the protocol in this document OR to use the protocol that is routinely used in their laboratory/institute/university. What matters is to specify as clearly as possible the method in the excel template (LakeName_ZooSize.xlsx), in the respective comment cells. To aid in further interpretation and homogenization of datasets, and if you are able to, the protocol should be shared with the ZooSize team champions.

Table of contents

. Some key pointsp2	
2. Measuring zooplanktonp3	,
Bosmina	
Calanoids	
Cyclopoids	
Daphnia	
Diaphanosoma	
Holopedium	
Leptodora	
Nauplii	
Others / rotifers	
S. Counting zooplankton - Links to protocolsp9	
1. EPA - Great Lakes	
2. Ontario - James Rusak	
3. Burrishoole LTER - Elvira deEyto	
4. DklT laboratory - Valerie McCarthy and Maria Caldero	
. Versions and updatesp1	0
5. Referencesp1	0

¹ ZooSize project description: https://rosalieb.github.io/rosaliebruelweb/ZooSize.html

² Thanks to Mireia Bartrons, Jessica Beyer, Sandra Brucet, Elvira deEyto, Jon Doubek, Fabio Lepori, Valerie McCarthy, and Jim Rusak, for sharing their protocol / answering our questions / submitting the first datasets that allowed us to get a sample of the diversity of methods!

³ Maria Caldero Pascual, Rosalie Bruel, Lauren Barth

1. Some key points

- We are asking for data measured by microscope in a first step, but the data call is also fitted for other methods (e.g., Flowcam, optical plankton counter), as there are many interesting questions we can answer with diverse sampling methods.
- Our goal is to obtain sample data with at least 200 total individual length measurements OR at least 20 individual length measurements per dominant taxa.
- We ask that you differentiate between copepodites and nauplii life stages (only when
 data are available or if a new campaign is carried out for this project: no need to go
 back to the sample if it has already been processed). Some data providers have also
 added rotifers and adult crustaceans' sex (i.e., male or female) and in case of
 females if they were carrying eggs (e.g., +eggs). Feel free to add this information as
 an extra column linked with each measurement provided, even though it is not
 required at this stage.
- Additionally, we ask that, if possible, you provide a photo identifying how each taxa
 was measured for length, so we are aware of potential protocol differences (e.g.,
 measurements including vs excluding helmets of *Daphnia*).

Guideline to identify different copepodite life stages (from P1 to P5 or adult)

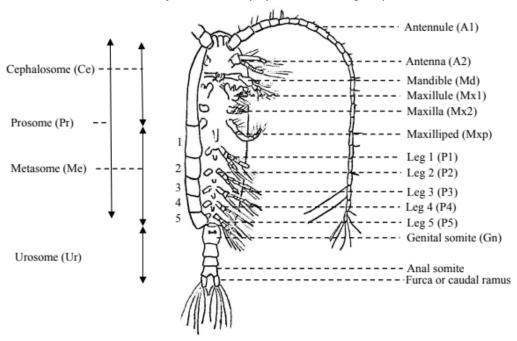


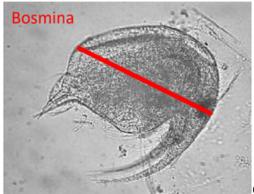
Fig. 1. Diagrammatic illustration of the external morphology and appendages of a CVI♀ calanoid copepod. Abbreviations for parts are shown and the pedigerous segments are numbered (Based on Mauchline, 1998).

2. Measuring zooplankton

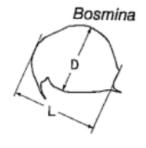
The table below summarises how to measure the different taxa. Refer to the following pages for examples and pictures.

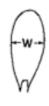
Taxa name	Recommended measurement (see photos below)	References
Bosmina sp.	Length from the top of the head to the end of the carapace.	Standard Operating Procedure for Zooplankton Analysis, Great Lakes, LG403 (2016)
Bythotrephes sp.	Body length, excluding the caudal process.	Standard Operating Procedure for Zooplankton Analysis, Great Lakes, LG403 (2016)
Calanoids	From top of the head to tip/extremity of caudal rami not including caudal setae	Malley et al. (1989) See measurement '*' in image by Bottrell et al (1976)
Cercopagis	Body length, from the top of the eye to the end of the caudal claws.	Standard Operating Procedure for Zooplankton Analysis, Great Lakes, LG403 (2016)
Cyclopoids	Length from the top of the head to the tip/extremity of caudal rami not including caudal setae	Malley et al. (1989) See measurement '*' in image by Bottrell et al (1976)
<i>Daphnia</i> sp.	Length from top of the head to anterior base of the spine not including the tail.	Malley et al. (1989)
<i>Daphnia</i> sp. with helmet	For the cyclomorphotic forms, if possible measure both from the anterior edge of the eye to the base of the tail spine (defined as standard length = SL in this study) and from the anterior margin of the helmet to the base of the tail spine (defined as total length = TL in this study).	Culver et al. (1985)
Diaphanosoma sp.	Length from the top of the head to the base of the caudal spine.	Malley et al. (1989)
Holopedium sp.	Length from the top of the head to the end of the carapace.	Malley et al. (1989)
Leptodora sp.	Length from the top of the head to the end of the carapace excluding spines.	Standard Operating Procedure for Zooplankton Analysis, Great Lakes, LG403 (2016)
Nauplii	Along the longest linear distance	SOP Lake Ontario shared by James Rusak

Bosmina sp.



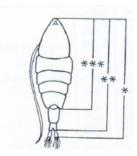
© Fabio Lepori



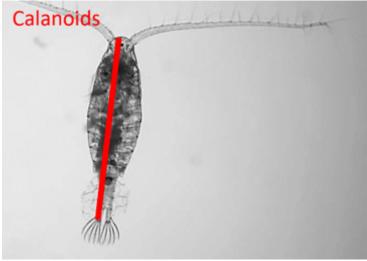


Malley et al. (1989)

Calanoids

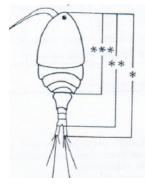


Bottrell et al (1976) - measure the length indicated by a single *

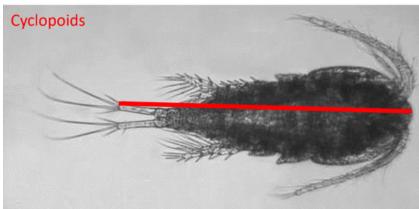


© Fabio Lepori

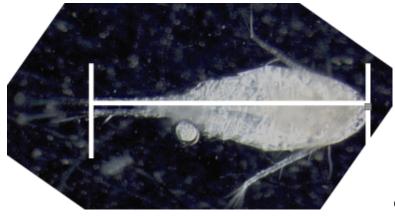
Cyclopoids



Bottrell et al (1976) - measure the length indicated by a single *

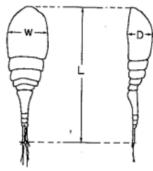


© Fabio Lepori



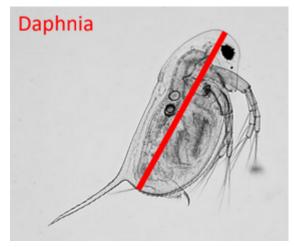
© James Rusak



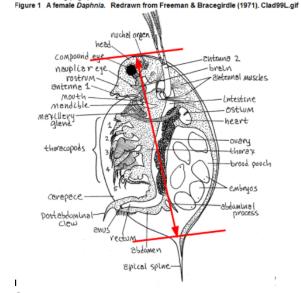


Malley et al. (1989)

Daphnia sp. (without helmet)



© Fabio Lepori

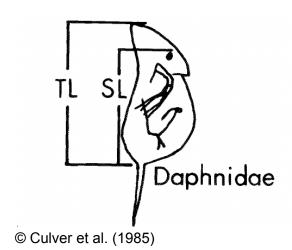


© Elvira de Eyto



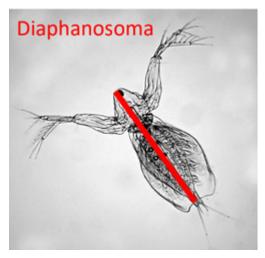
© James Rusak

Daphnia sp. with helmet

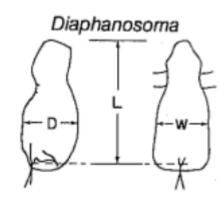


6/10

Diaphanosoma sp.

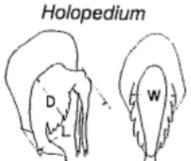


© Fabio Lepori



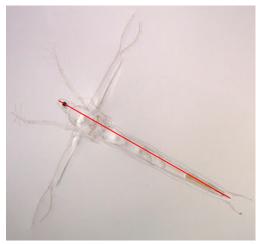
Malley et al. (1989)

Holopedium

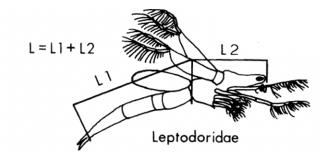


Malley et al. (1989)

Leptodora sp.



© Mireia Bartrons



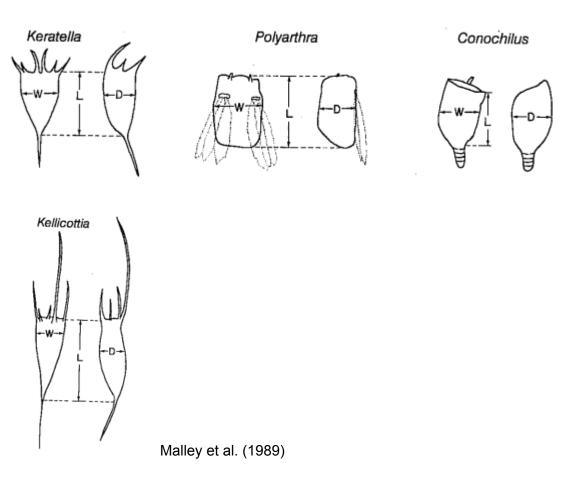
© Culiver et al. (1985)

Nauplii

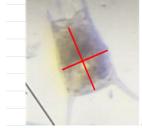


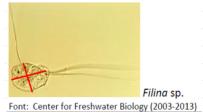
Naupli © Mireia Bartrons

Others / rotifers









Filina sp.

Keratella sp. I

Keratella sp. II

© Mireia Bartrons

3. Counting zooplankton - Links to protocols

Below are links to 4 protocols displaying a range in complexity and details. They all provide ways of estimating abundances from zooplankton samples. We accept data collected with any of these protocols. If you already have a protocol within your lab, and you would rather keep using it, it isn't a problem; just document your methods precisely in the data form.

Note on preservatives - we are working on this question and may come up with an updated SOP once we gather enough feedback.

The preservative used influences the real zooplankton dimensions (Jaspers and Carstensen, 2009, https://doi.org/10.4319/lom.2009.7.430). For example, the use of formalin shrinks zooplankton while the use of ethanol bloats animals. There are existing published correction factors to account for this (e.g., Jaspers and Carstensen, 2009). We therefore will need to know your protocol in detail to correct for these differences.

In the meantime, we advise to narcotise the animals within one hour of collection with soda water/carbonate water (e.g., <u>Great Lakes SOP</u>⁴; Ordonez, 2010; Culver, 1985) or Alka-Seltzer® tablet (Maria Caldero and Jon Doubek use that method for instance), and then use sucrose formalin solution*.

* Sucrose formalin solution

REAGENTS

- Sucrose (crystalline)
- Formalin (37% solution of formaldehyde in water)
- Borax (powder)

BUFFERED SUCROSE FORMALIN PREPARATION

- 1. Dissolve 60 grams of sucrose in 1000 mL of formalin (Haney and Hall, 1973)
- 2. Then dissolve 9 grams of borax in 1000mL of sucrose formalin
- 3. Store in a labeled plastic container
- 1. EPA Great Lakes
- 2. Ontario, Zebra2-Dorset method James Rusak shared protocol
- 3. Lough Feeagh Long-Term Monitoring Program Elvira deEyto
- 4. Valerie McCarthy and Maria Caldero protocol

⁴ Great Lakes SOP - specifics for sample collection and preservation steps: "The zooplankton samples should be refrigerated as soon as possible after collection. In the shipboard biology lab, 20 mL of soda water is measured with a graduated cylinder, into the sample to narcotize the organisms within 1 hour of sample collection. The sample then stands for 30 minutes in the refrigerator. Under a hood, 20 mL of sucrose formalin solution is added to the sample. The sample storage bottle (500-mL plastic sample bottles) is filled to the top with reagent water and tightly capped, the cap and neck are wrapped with parafilm to prevent leaks, and the sample storage bottle is stowed in a designated cooler in the walk-in refrigerator."

4. Versions and updates

This is the first version (v1.0) circulated within the group.

We will be working on adding comments on the use of preservatives - lookout for updates (either consult the <u>project's page</u> OR email us).

5. References

Some collected pdfs:

https://drive.google.com/drive/folders/1Gq--2ztR35Nv9vstT9oRQaWMJrdRXw5M?usp=sharing

Papers regarding correcting factors depending on the preservative used:

https://www.tandfonline.com/doi/abs/10.1080/17451000.2021.1900576

https://academic.oup.com/plankt/article/18/4/483/1490972?login=true

https://academic.oup.com/plankt/article-abstract/16/12/1601/1461253

https://www.scielo.br/j/ni/a/RbpB6YdFMLZB3tcgx6vVjfL/?lang=en