



INTRODUCTION

Starvation has a major impact on physiological processes in marine plankton and, in turn, these processes have a major impact on the energy transfer through marine ecosystems. How starvation changes different aspects of the energy transfer has been understudied because planktologists have largely relied on biomass fractionation as a research tool. Here, with a caridean shrimp, *Palaemon elegans*, we experiment with the cellular energy allocation approach (CEA index), a new index that is calculated as the ratio between energy available (E_a), to energy consumption (E_c). It allows us to evaluate the net energy budget (Verslycke and Janssen, 2002). We analyzed the effect of 72 hours of starvation on the CEA index, the respiratory electron transport system activity (ETS), and on the shrimps' lipids (LIP), carbohydrates (CARB) and proteins (PROT) to help further our understanding on the energy transfer in the shrimp. The E_a equals the sum of the LIP, CARB, and PROT. The E_c equals the ETS activity in comparable units.

METHODOLOGY

CARB

- Dubois et al. (1956)
- 150 μ L sample + 150 μ L phenol (5%) + 750 μ L H₂SO₄
 - 10min, vortex
 - Incubation 30°C, 10min
 - Read absorbance at 485nm (Standard: Glucose in buffer)

LIP

- Extraction (1:1:0.9; Bligh and Dyer, 1959)
 - 250 μ L sample + 500 μ L chloroform + 500 μ L methanol + 200 μ L water
 - 500 μ L standard + 500 μ L methanol + 450 μ L water
 - Centrifugation (2600g, 10min, 4°C)
- Charring (200°C, 15min): 100 μ L organic phase + 500 μ L H₂SO₄
- Colorimetric assay (Barnes and Blackstock, 1973):
 - 40 μ L charred sample + 1mL phosphovanillin reagent
 - Incubation 37°C, 15min
 - Read absorbance at 525nm (Standard: Oleic acid in methanol)

PROT

- Smith et al. (1985)
- 250 μ L sample + 250 μ L work reagent
 - Incubation 37°C, 30min
 - Read absorbance at 540nm (Standard: BSA in buffer)

SAMPLE PRE-TREATMENT

- Homogenization in 0.1M sodium-potassium phosphate buffer
- Centrifugation: 4000rpm, 10min, 0°C



ENERGY BUDGET

$$\text{CEA index} = E_a / E_c$$

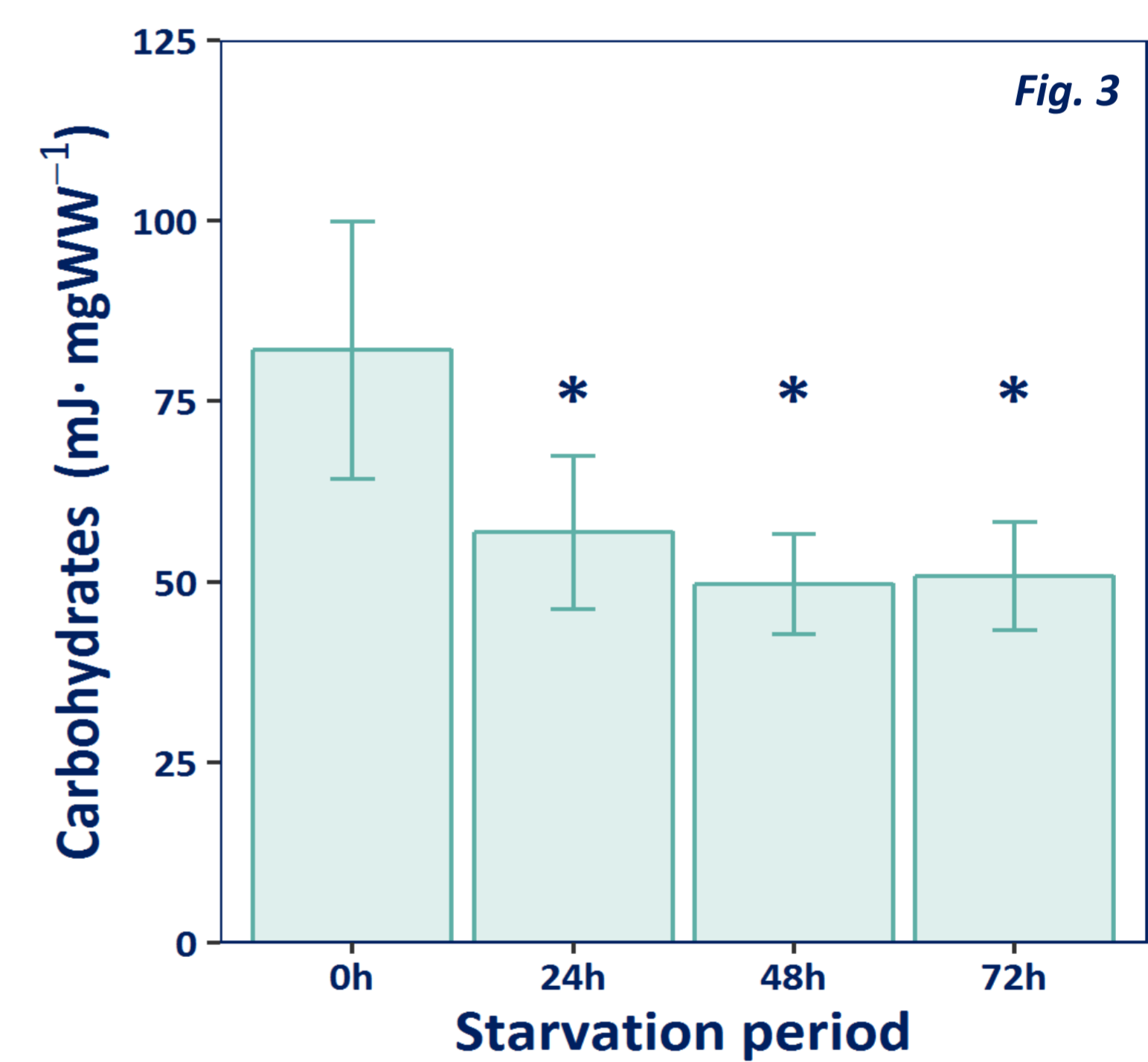
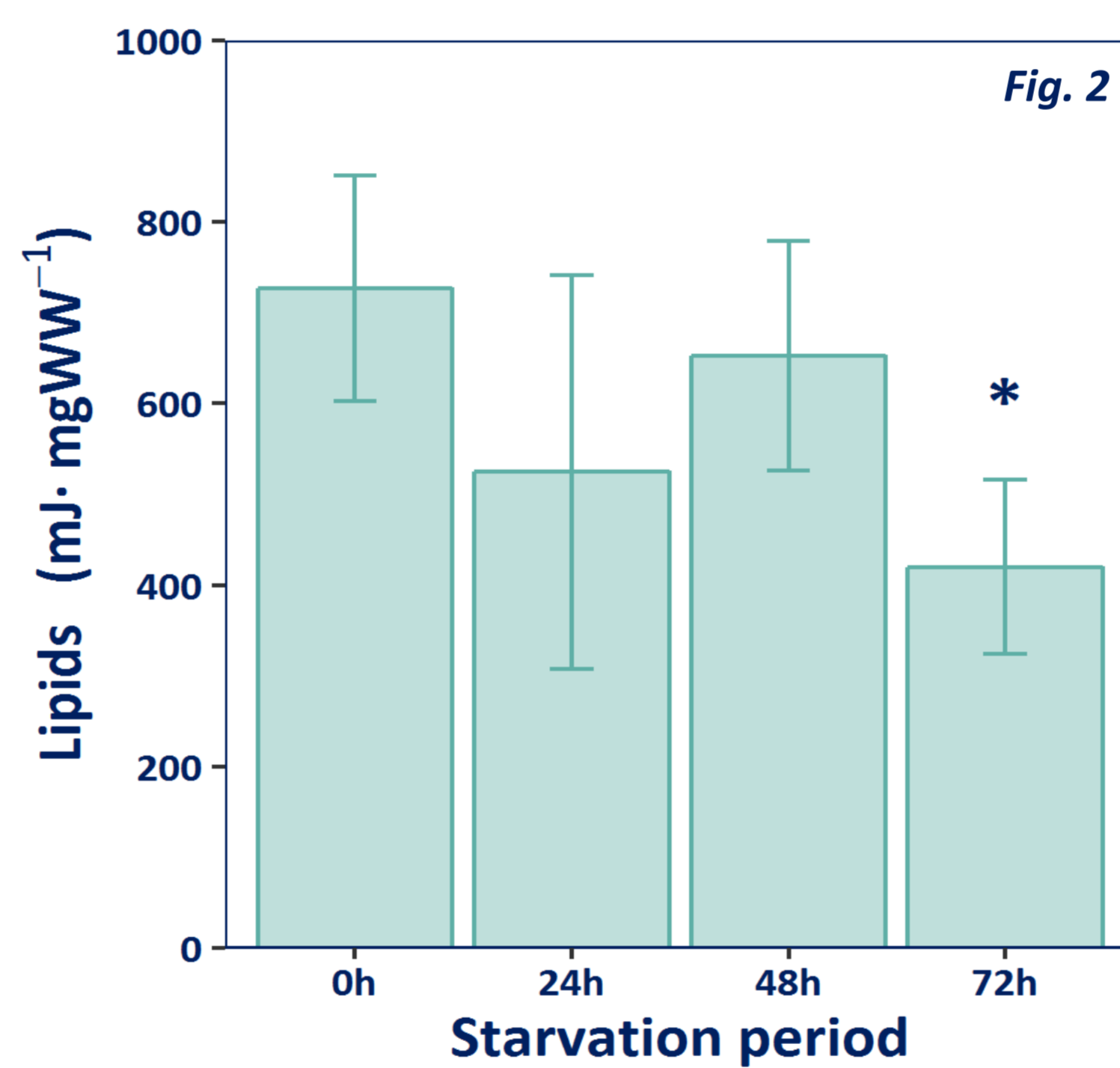
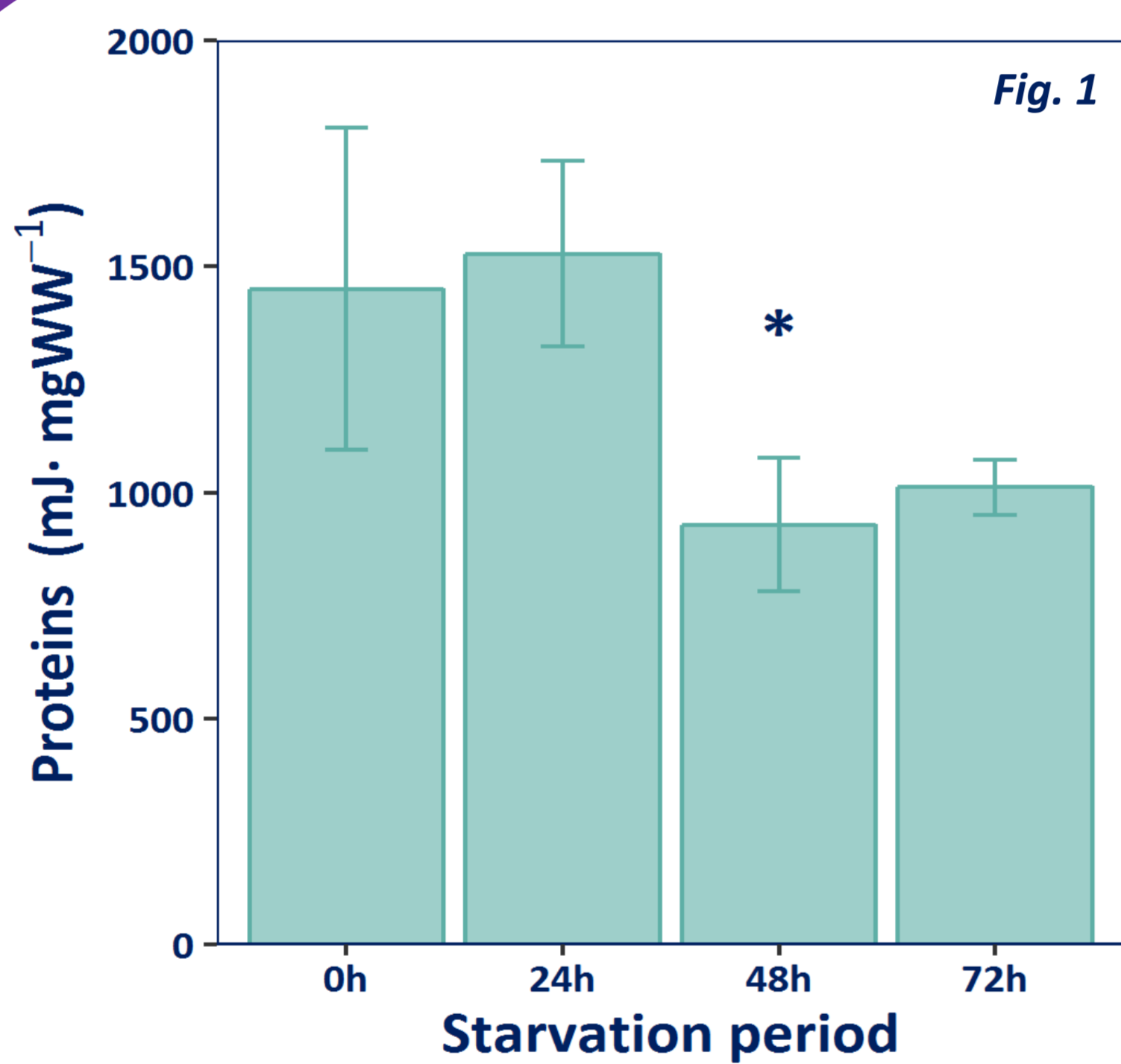
Parameters transformed into energy equivalents (Gnaiger, 1983)

ENERGY CONSUMPTION (E_c)

ETS

- Owens and King (1975); Gómez et al. (1996)
- 100 μ L sample + 300 μ L substrate solution + 100 μ L INT
 - Read absorbance kinetically at 490nm during 8 min

RESULTS

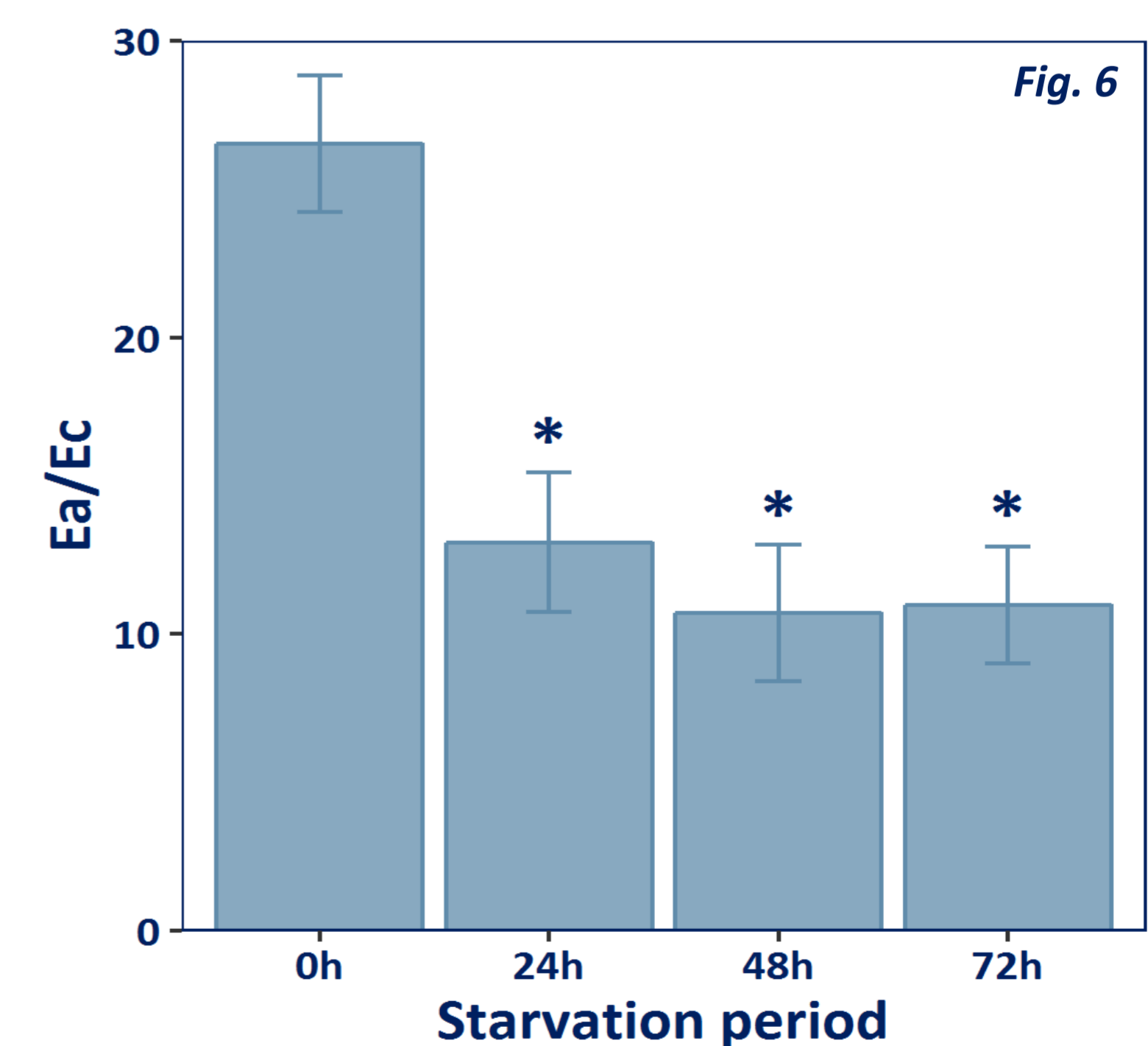
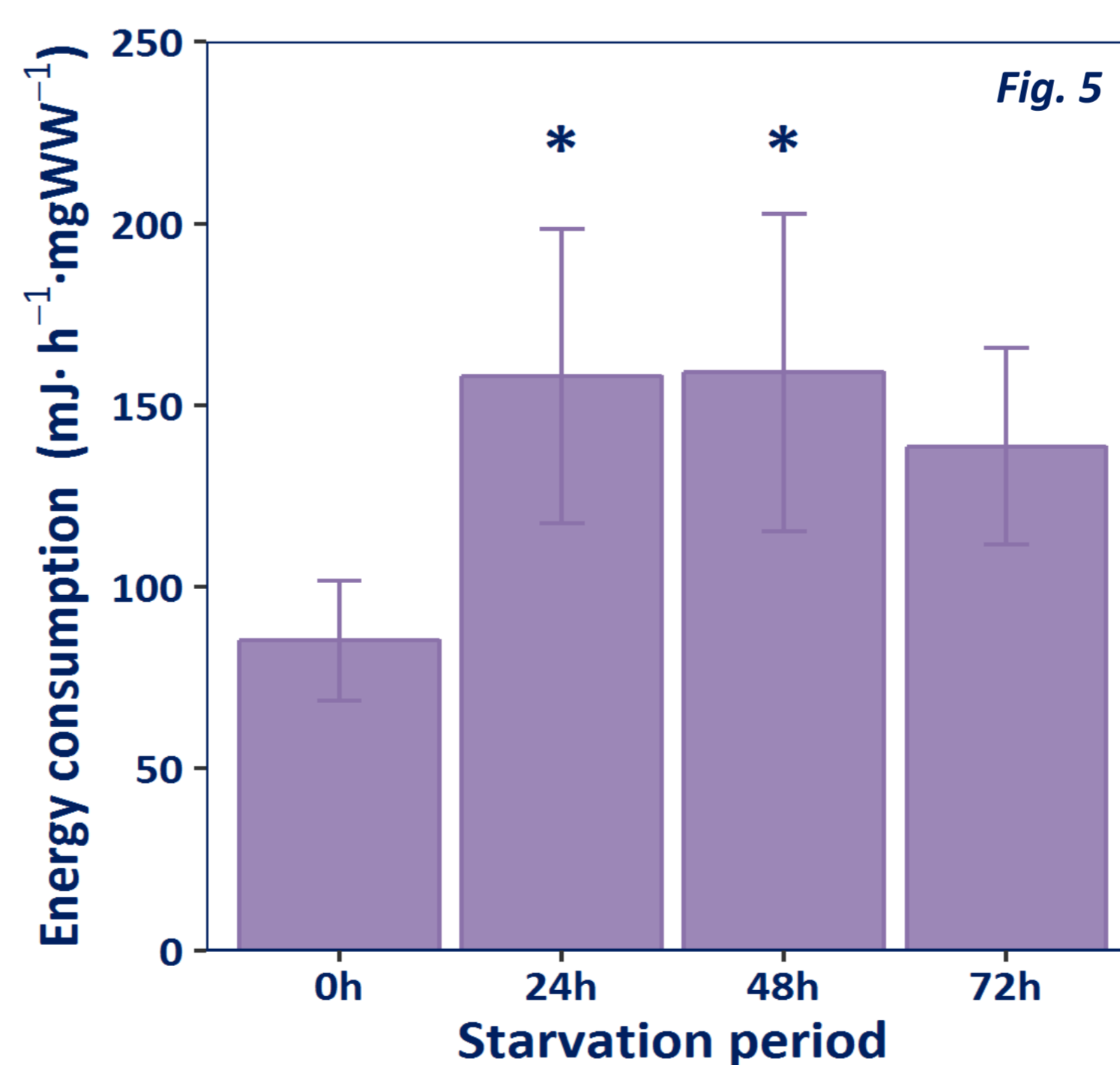
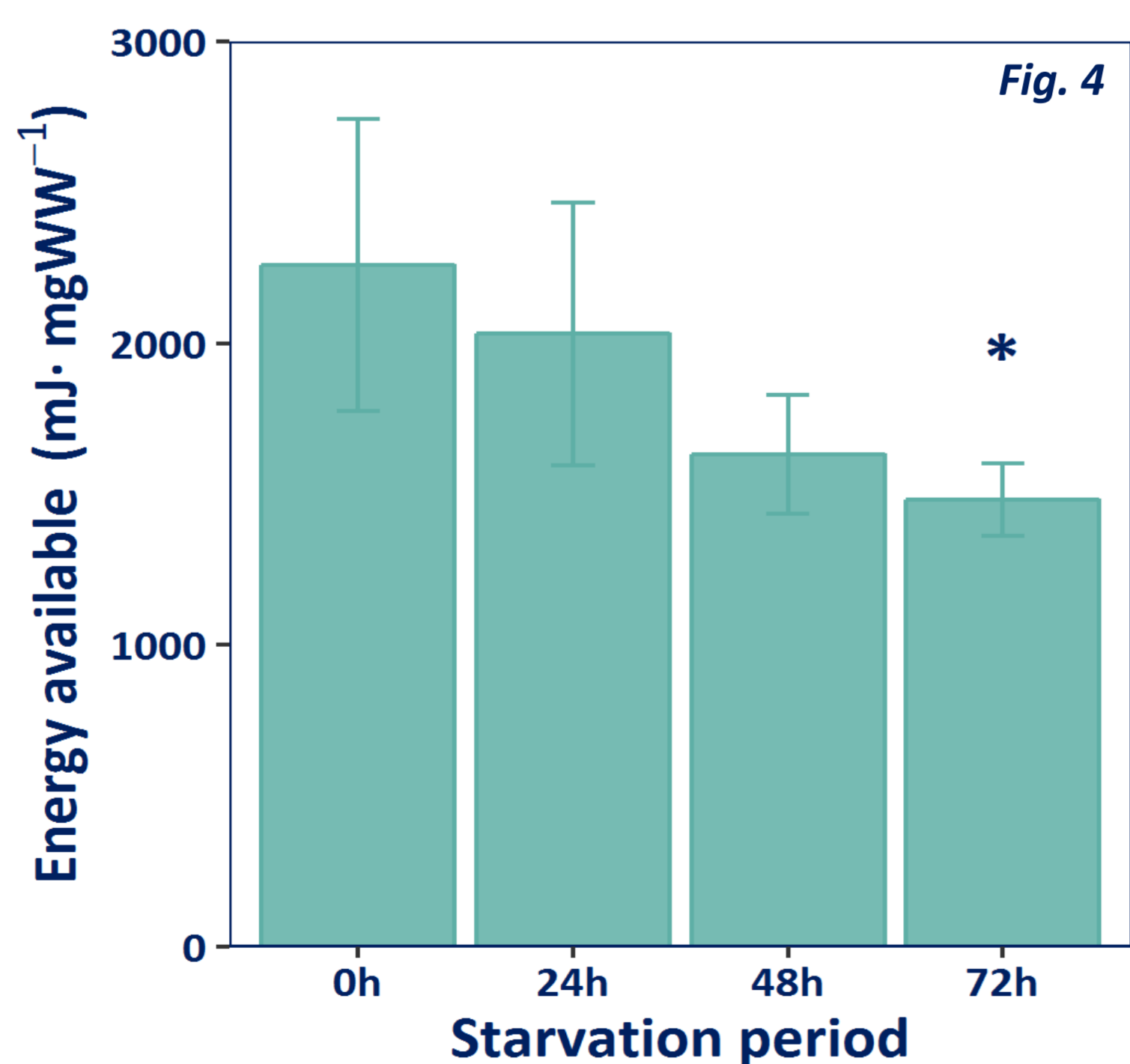


(*) Significant differences compared with the control (0h)

- After 24h: Carbohydrates dropped by 38.2% (Fig. 3)
- After 48h: Proteins had decreased by 36.0% (Fig. 1)
- After 72h: Lipids had dropped by 42.3% (Fig. 2)

- Energy available: significant decreasing trend of 34.42% at 72 h. (Fig. 4)
- Energy consumption: significant increase at 24h that stabilized after 72h. (Fig. 5)

ENERGY BUDGET (Fig. 6)
Energy available was higher than the energy consumption during the experiment. The ratio (E_a/E_c) decreased for the first 24h then remained stable up to 72h.



CONCLUSIONS

- ❖ Proteins were the main energy reserve (1450.8 ± 355.7 mJ·mgWW⁻¹) followed by lipids (727.2 ± 121.4 mJ·mgWW⁻¹) and carbohydrates (82.1 ± 17.9 mJ·mgWW⁻¹).
- ❖ Carbohydrates were the first energy source mobilised (after 24h), followed by proteins (48h), and lipids the last (at 72h).
- ❖ Energy regulation shows a rapid shift mechanism of *P. elegans* under adverse conditions.

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