

Characteristics of Iridophore Crystals in *Anolis carolinensis*Jiahui Hu¹, Xu Chen^{1,*}, Aiping Liang^{1,2,*}¹ Tianjin Key Laboratory of Conservation and Utilization of Animal Diversity, College of Life Sciences, Tianjin Normal University, Tianjin, China² Institute of Zoology, Chinese Academy of Sciences, Beijing, China

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INTRODUCTION & AIM

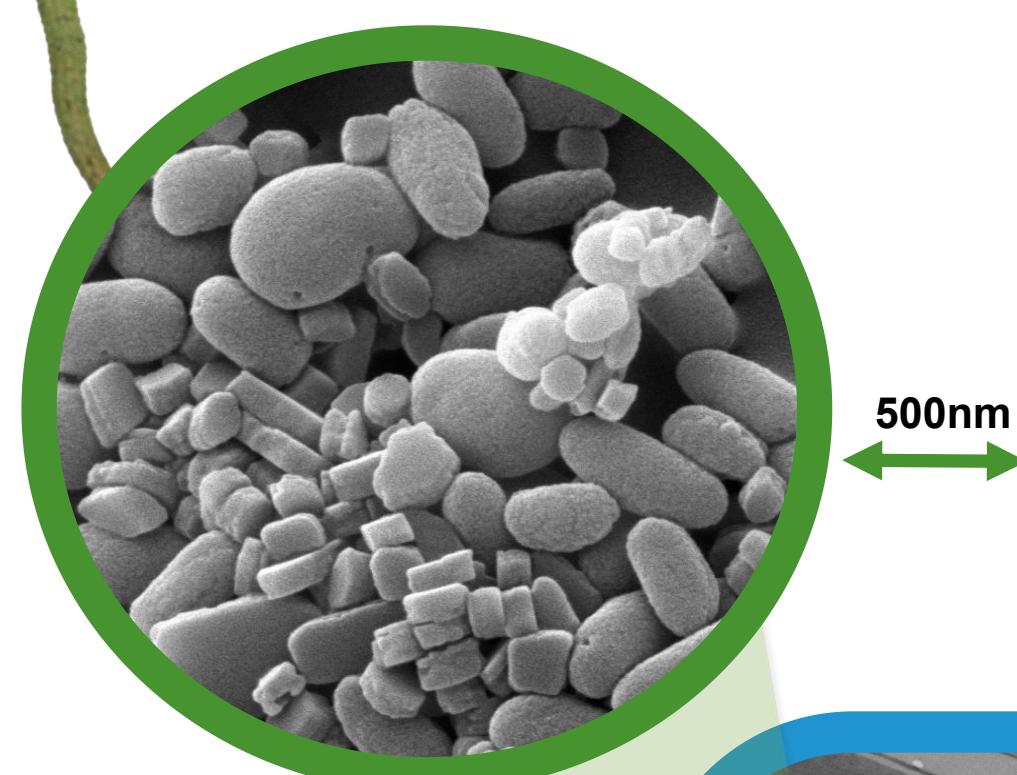
Anolis carolinensis

INTRODUCTION

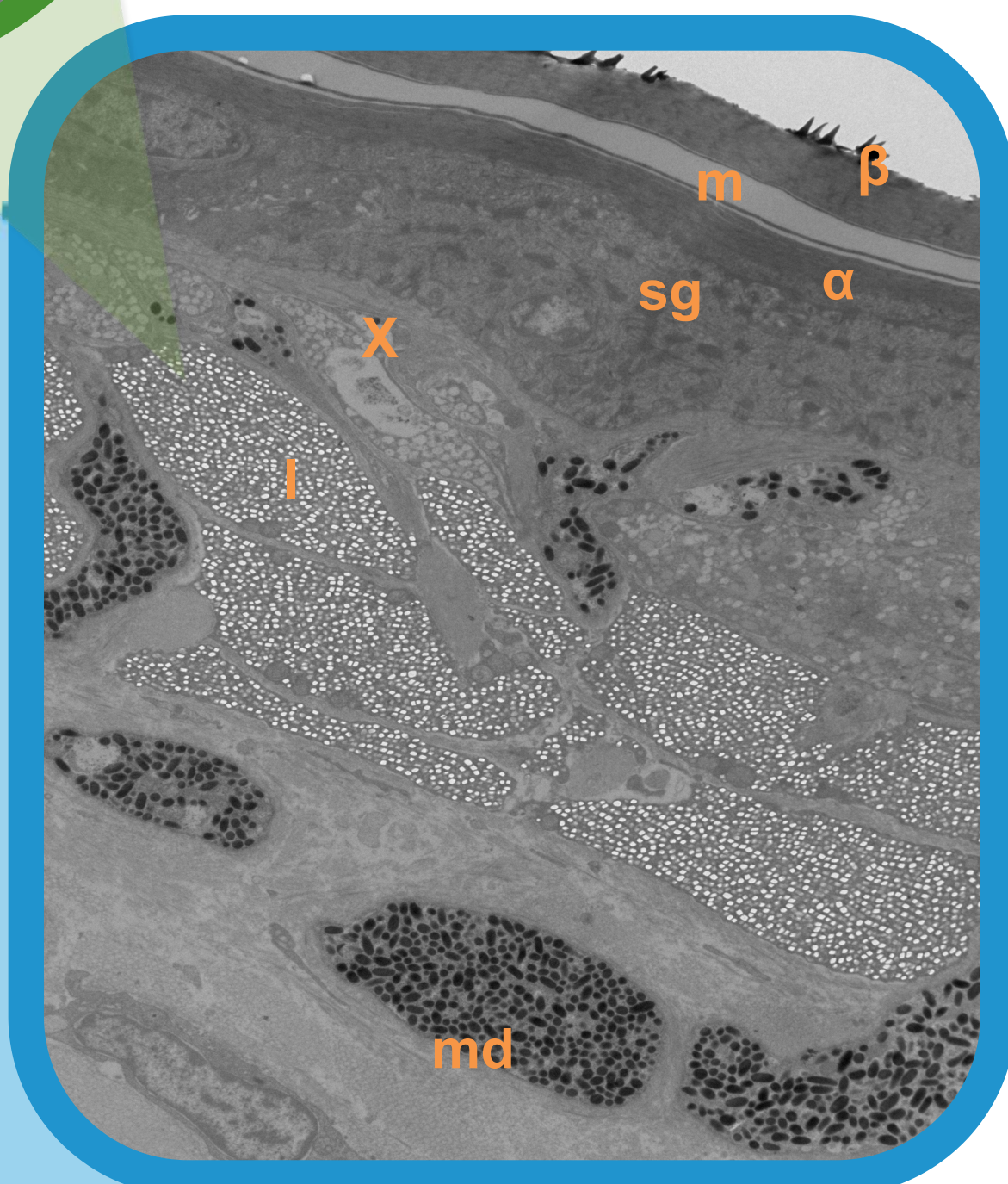
Chameleons or lizards are biomimetic organisms commonly used for color-changing or thermostatic materials. However, the structure of purine crystals within iris cells and their specific role in body color regulation are still poorly understood.

This study aims to address the aforementioned knowledge gap by selecting *Anolis carolinensis* (a lizard species with the ability of rapid body color change) as the experimental subject, with a focus on investigating the structure of purine crystals in its iridophore cells and clarifying their specific function in body color regulation.

Three environmental temperatures (20°C, 30°C, 40°C), all within the range that ensures the normal physiological activity of lizards, were selected to investigate the relationship between environmental temperature and skin reflectance (300–700 nm). →

Transmission
electron microscopy

α : α -keratin layer
 β : β -keratin layer
 m : intermediate differentiation layer
 sg : stratum germinativum
 x : xanthophore
 i : iridophore
 md : melanophores

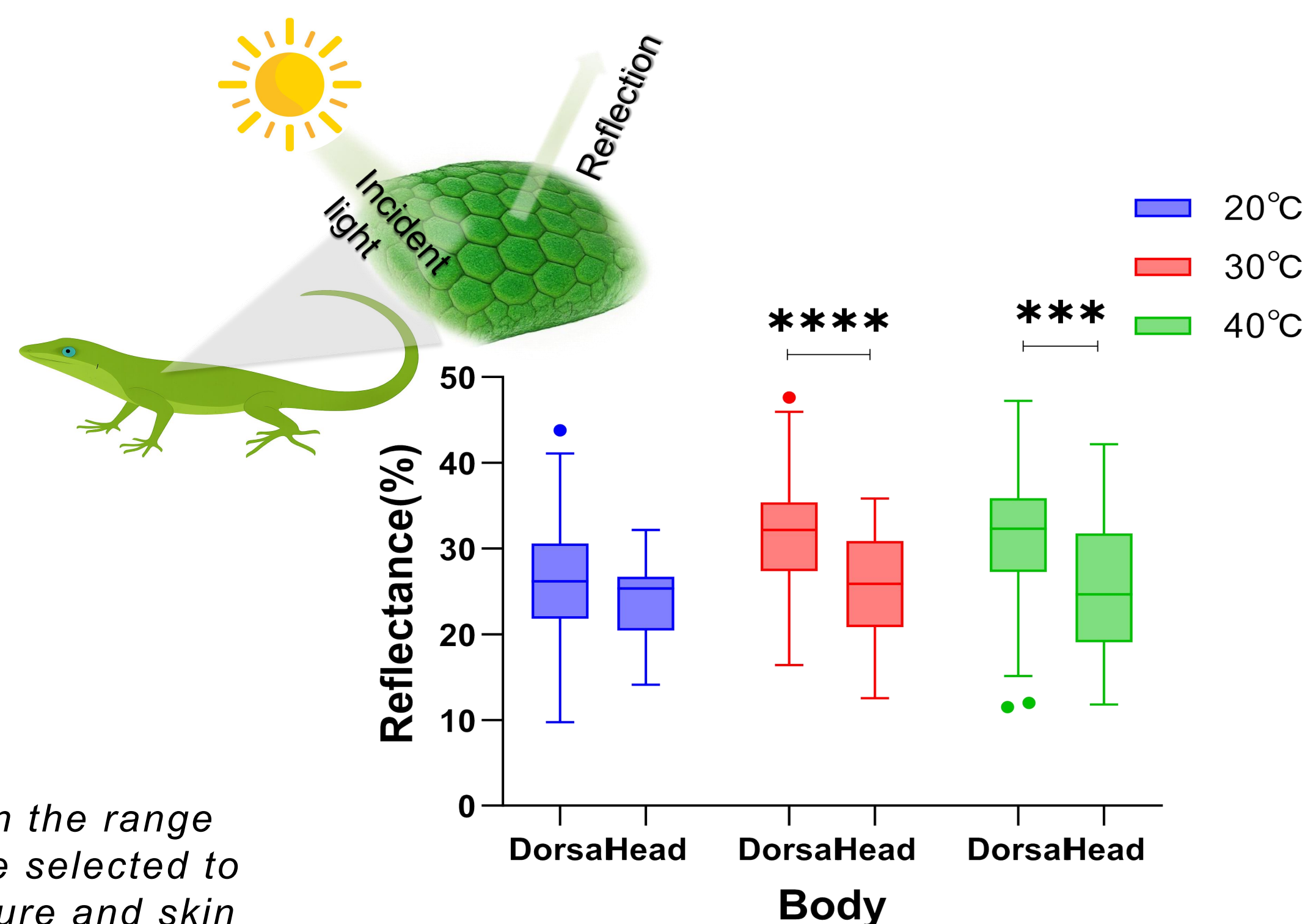


REFERENCES

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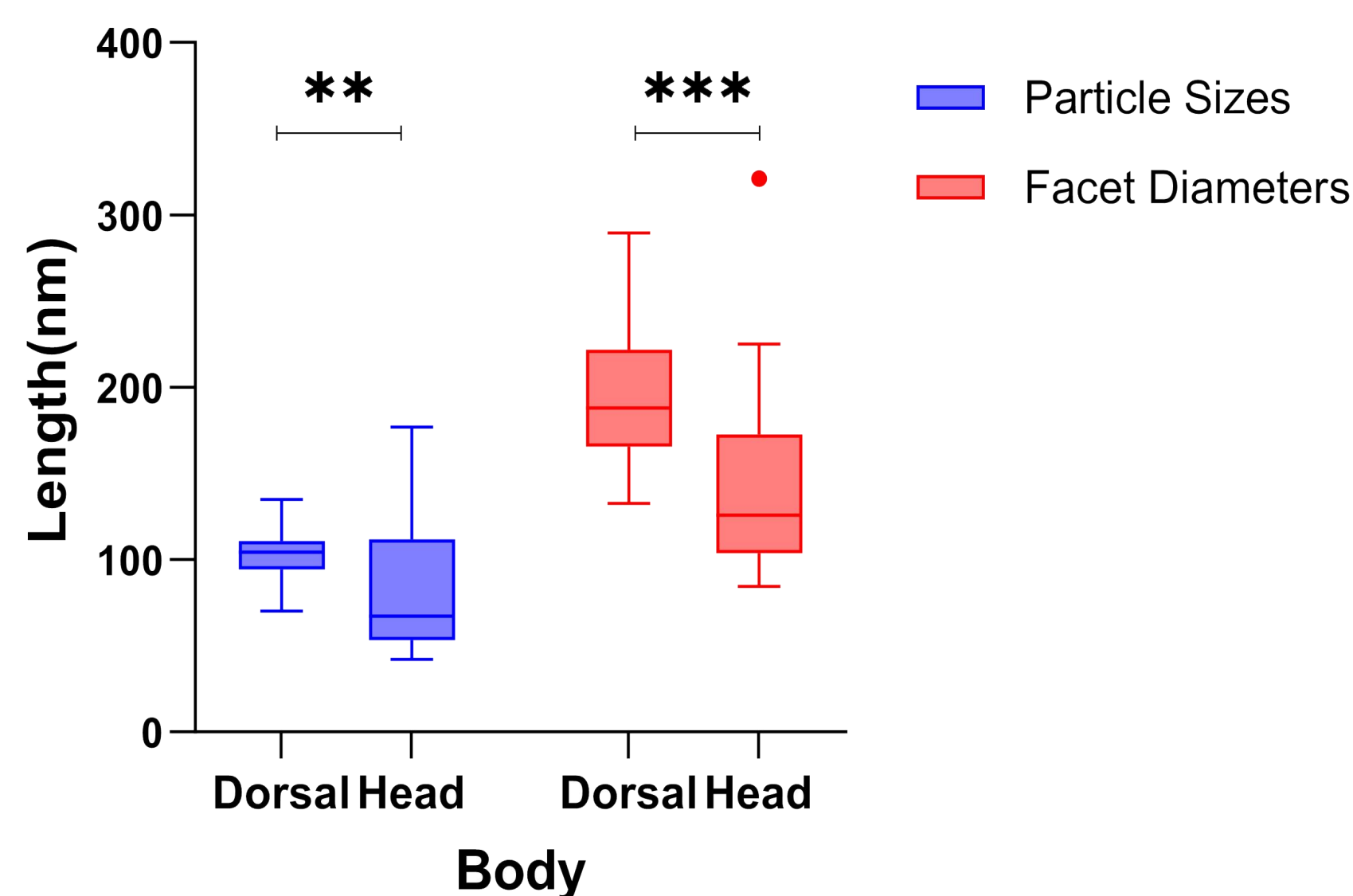
RESULTS & DISCUSSION



Skin reflectance of different body parts at different ambient temperatures

At different ambient temperatures, the peak wavelength of reflection increases with the increase of temperature, and there is a significant difference between the head and dorsal. The dorsal thoracic peak increased from $26.02 \pm 0.58\%$ at 20 °C to $31.40 \pm 0.66\%$ at 40 °C. The peak of the head increased from $23.97 \pm 0.82\%$ to $25.52 \pm 1.41\%$.

Analysis of the differences in crystal size at different parts



Further examination of crystals extracted from iris cells using SEM revealed lumpy or short columnar structures that were often stacked on top of each other, with facet diameters of 172.46 ± 29.67 nm and particle sizes of 95.30 ± 13.13 nm. Significant differences in crystal size were observed between the head and dorsum, with the dorsolateral lens being thicker and the facet diameter larger than the crystals of the head.