

Effets de la castration sur l'immunoexpression de la matrilysine (MMP-7) dans les vésicules séminales du Mérion de Libye (*Meriones libycus*)Mansouria Belhocine<sup>1,2</sup>, Ramadhan Hatti<sup>1,2</sup>, Thérèse Gernigon-Spychalowicz<sup>2</sup>1: Laboratory of Beneficial Microorganisms, Functional Foods and Health (LMBAFS), FSNV, Abdelhamid Ibn Badis University of Mostaganem, Algeria. [manbelhocine@hotmail.com](mailto:manbelhocine@hotmail.com)

2: Laboratory of Arid Areas Research (LRZA), Reproduction of Small Vertebrates, Faculty of Biological Sciences (FSB), USTHB, El Alia, Algiers, Algeria

## INTRODUCTION

MMPs are involved in physiological and pathophysiological tissue remodeling. They play important roles in reproduction from development to adulthood. We studied, immunohistochemically the MMP-7 in order to demonstrate its involvement in the physiology and seasonal activities of *Meriones libycus* seminal vesicles.

## MATERIALS AND METHODS

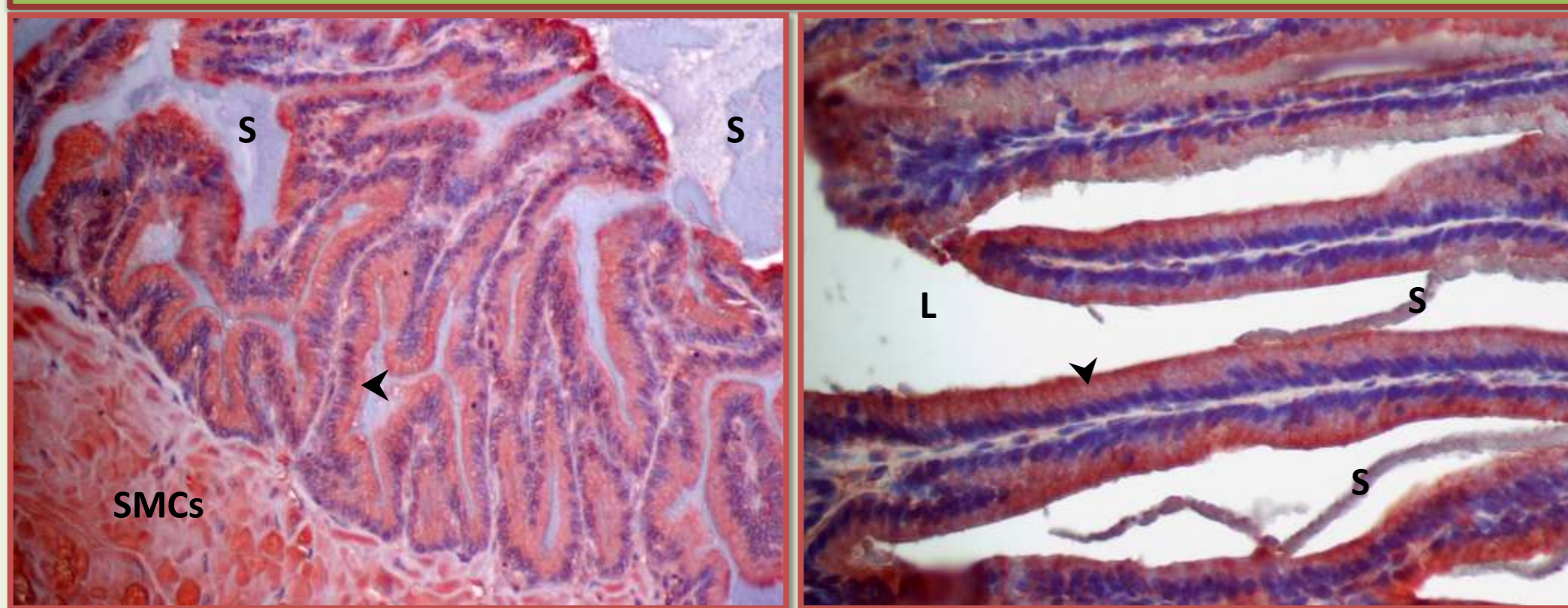
The Libyan jird was collected in the breeding period and the resting season from Béni-Abbès area, in the Algerian Sahara. The seminal vesicle, also taken from castrated Libyan jird for 3, 7, 30, 50 and 90 days in spring, was submitted to the indirect immunohistochemical protocol with amplification with streptavidin-biotin-peroxidase and AEC/DAB as chromogen.

## ANIMALS

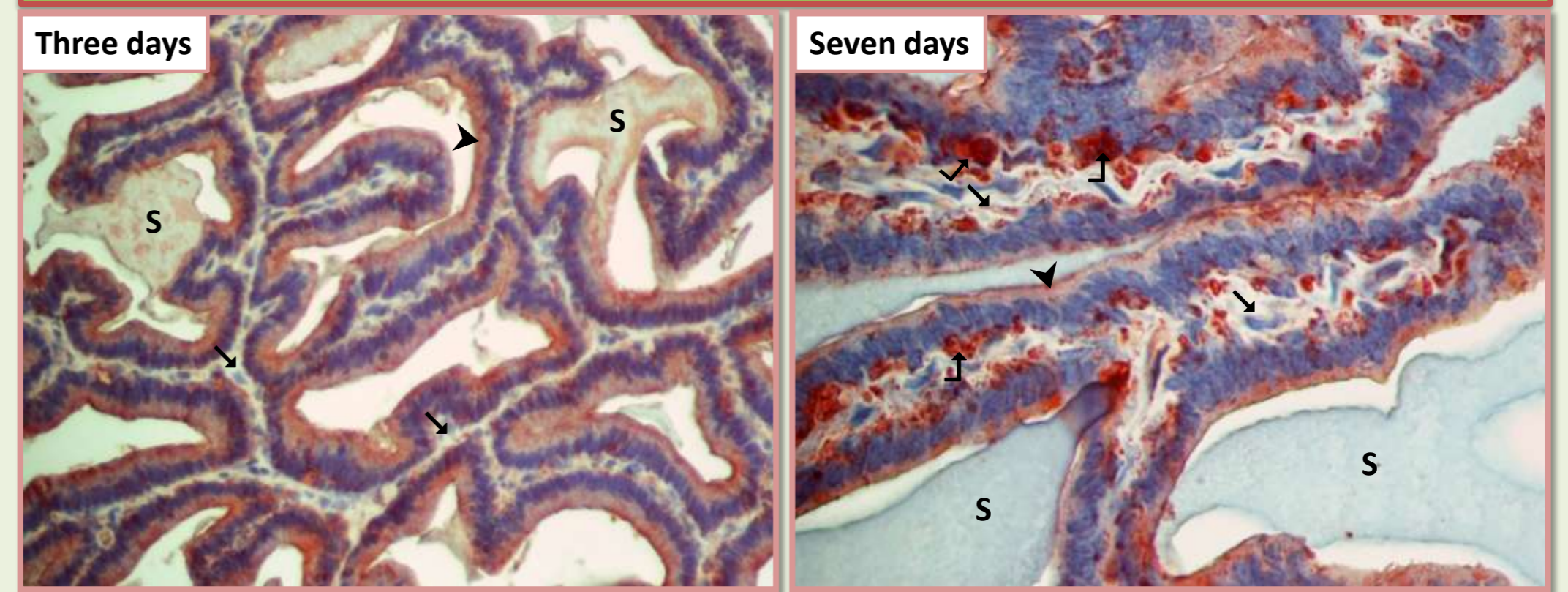
*Meriones libycus* is a nocturnal herbivorous and granivorous Saharan Rodent belonging to the Gerbillidae family. It lives in a superficial burrow arranged under the most important bushes. So, it benefits from the shade procured by the plants (Petter, 1961).

- **Breeding period:** spring and early summer.
- **Resting phase:** late summer, autumn, late winter.

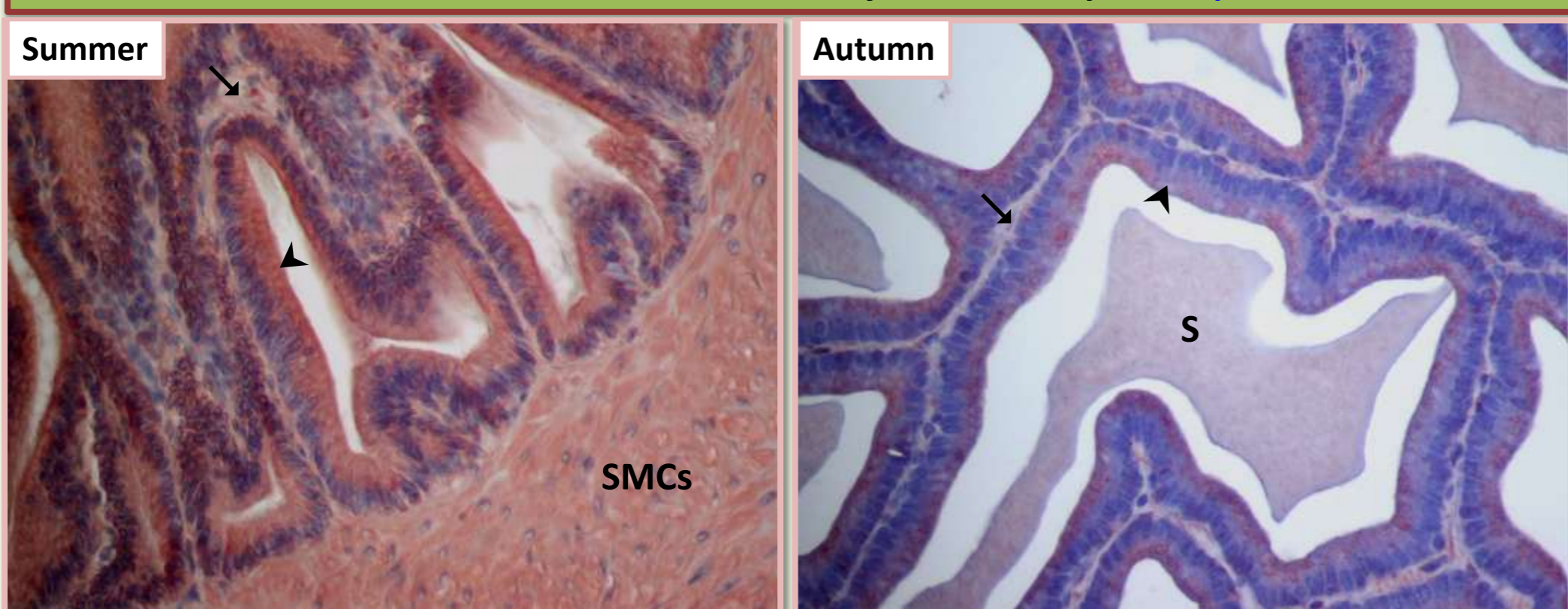
## RESULTS

MMP-7 immunoreponse in the seminal vesicle of *Meriones libycus* in the breeding period

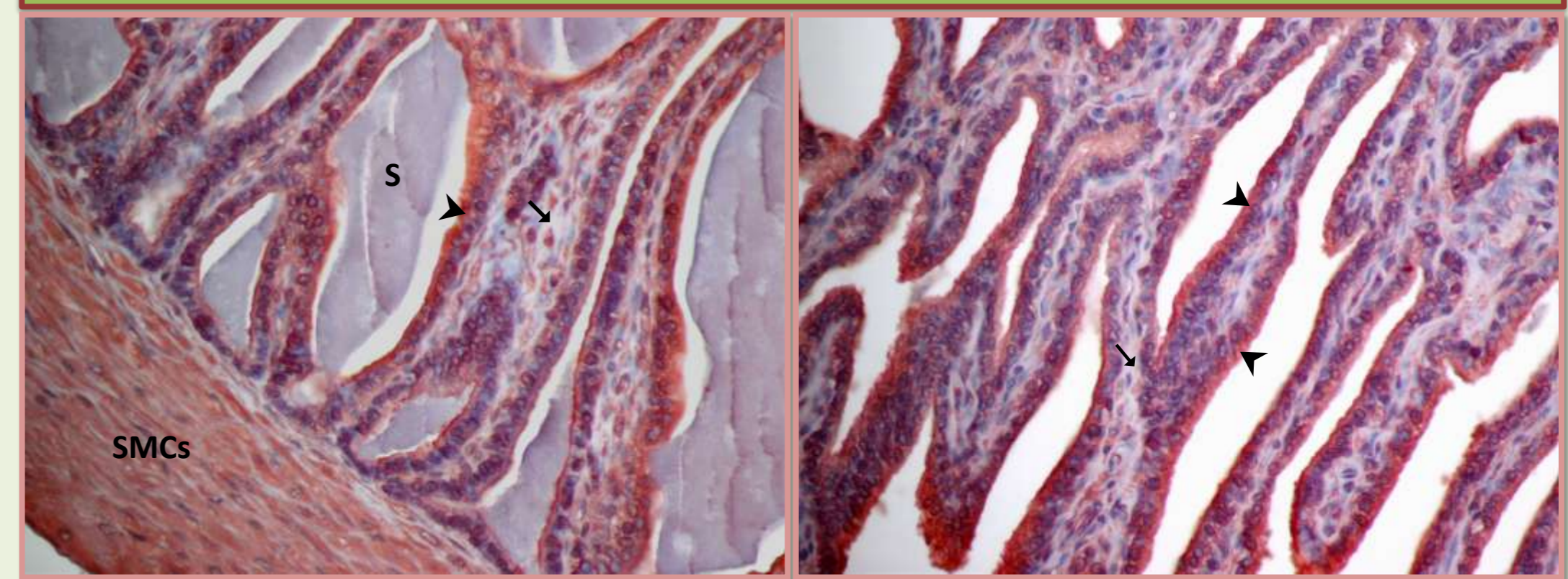
A non immunostained secretion (S) is observed in the Lumen (L) of the seminal vesicle. Luminal epithelial cells (▶) and smooth muscle cells (SMCs) show a strong immunohistochemical staining. No immunoreactivity in the extracellular matrix (↘).

MMP-7 immunoeexpression in the seminal vesicle of a short period castrated *Meriones libycus*

A weak positive immunohistochemical staining is seen in the luminal epithelial cells (▶). The extracellular matrix (↘) and the secretion (S) show a negative immunoreactivity. In seven days of castration a significant peri-basal (J) immunoreponse is observed.

MMP-7 immunodetection in the seminal vesicle of *Meriones libycus* in quiescence season

Luminal epithelial cells (▶) with an attenuated immunolabeling in autumn. The extracellular matrix (↘) and the secretion (S) are not immunostained.

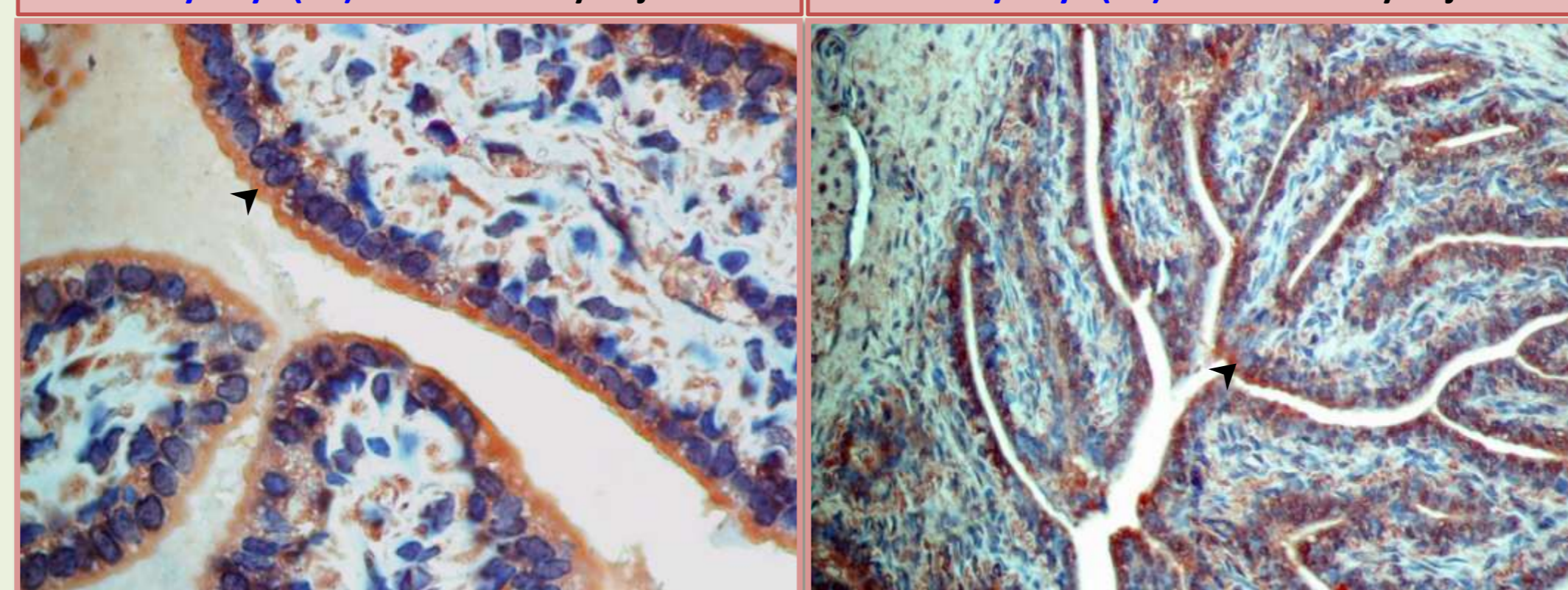
MMP-7 immunostaining in the seminal vesicle of thirty days castrated *Meriones libycus*

The pattern of the immunostaining is the same as that of the quiescence period in summer: Luminal epithelial cells (▶) and smooth muscle cells (SMCs) show an important immunoreactivity. The extracellular matrix (↘) and the secretion (S) are not immunostained.

MMP-7 immunoreaction in the seminal vesicle of a long period castrated *Meriones libycus*

## Fifty days (50) castrated Libyan jird

## Ninety days (90) castrated Libyan jird



The MMP-7 immunostaining is concentrated in the luminal epithelial cells (▶)

## DISCUSSION AND CONCLUSION

In the **breeding period**, the MMP-7 is highly expressed in luminal epithelial cells and is absent in the extracellular matrix (ECM) and the secretion. This immunoreaction decreases in the **three days castrated** *Meriones libycus* and in **seven days castrated** *Meriones libycus* the immunostaining is delocalized in the peri-basal ECM suggesting a role of MMP-7 in epithelial atrophy induced by hormonal deprivation. In the **quiescence season**, MMP-7 immunoreponse decreases slightly in epithelial cells; SMCs also show a net immunoreaction. This pattern of the immunolabeling remains as it is after **thirty days** of castration. However, following a **prolonged castration (50 and 90 days)** the immunoreactivity diminishes strongly in the luminal epithelial cells. Similar results were obtained in the canine testis, epididymis and semen (Warinrak *et al.*, 2015). These results indicate a seasonal fluctuation in the expression of the MMP-7 and a dynamic tissue distribution. This allows us to stipulate a modulating effect of testosterone but not alone on the MMP-7 and the involvement of this enzyme in the physiology of the Libyan jird seminal vesicles, in its atrophy processes and its seasonal tissue remodeling as in hamster ovaries (Shahed *et al.*, 2015). Other authors postulated that MMPs could have a function in fertilization as has been shown in mammals (Ferrer *et al.*, 2012) and *Xenopus* (Iwao *et al.*, 2014; Sato *et al.*, 2015).

## REFERENCES

- ✓ Ferrer M, Rodríguez H, Zara L, Yu Y, Xu W, Oko R. 2012. MMP-2 and acrosin are major proteinases associated with the inner acrosomal membrane and may cooperate in sperm penetration of the zona pellucida during fertilization. Cell Tissue Res 349(3), 881-895.
- ✓ Iwao Y, Shiga K, Shiroshita A, Yoshikawa T, Sakiie M, Ueno T, Ueno S, Ijiri TW, Sato K. 2014. The need of MMP-2 on the sperm surface for *Xenopus* fertilization: its role in a fast electrical block to polyspermy. Asian-Australas J Anim Sci 27(11), 1526-1531.
- ✓ Petter F. 1961. Répartition géographique et écologique des rongeurs désertiques de la région paléarctique. Mammalia 25, Suppl, 222 p.
- ✓ Sato K. 2015. Transmembrane signal transduction in oocyte maturation and fertilization: focusing on *Xenopus laevis* as a model animal. Int J Mol Sci 16(1), 114-134.
- ✓ Shahed A, Simmons JJ, Featherstone SL, Young KA. 2015. Matrix metalloproteinase inhibition influences aspects of photoperiod stimulated ovarian recrudescence in Siberian hamsters. Gen Comp Endocrinol 216, 46 - 53.
- ✓ Warinrak C, Wu JT, Hsu WL, Liao JW, Chang SC, Cheng FP. 2015. Expression of matrix metalloproteinases (MMP-2, MMP-9) and their inhibitors (TIMP-1, TIMP-2) in canine testis, epididymis and semen. Reprod Domest Anim 50(1), 48 - 57.