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LOW TESTOSTERONE LEVELS INDUCE APOPTOSIS VIA ACTIVE 3-CASPASE DEPENDENT SIGNALING IN THE BLADDER WALL OF MALE RATS

Hypothesis / aims of study

The bladder can be considered a target organ for testosterone action, and low testosterone levels possibly cause damage to bladder cells. Decreases in circulating testosterone due to aging have been associated with bladder dysfunction. Senile and orchiectomized rats have been used as animal models for studying hypoandrogenism. We set out to study whether hypoandrogenism influences bladder wall cell damage in castrated and senile male rats.

Study design, materials and methods
Thirty male Wistar rats were divided into three groups of 10 animals each: group I (3-months-old), sham animals; group II (27months-old), senile animals; group III (3-months-old), subjected to bilateral orchiectomy, and sacrificed eight weeks after the procedure. The bladders were rapidly excised, weighed and sent for analysis. Stereological assays on collagen fibers and immunohistochemical analysis with active 3-caspase were performed on the bladder cells.

Interpretation of results and conclusion

Bladder weights were greater in the senile group than in the other groups. Stereological collagen fiber analysis demonstrated higher density in group III than in groups I and II (p<0.05). The absolute density was 4.15 mm³ in group I, 22.3 mm³ in group II and 19.3 mm³ in group III. Semiquantitative active 3-caspase analysis showed greater percentages in the senile group II than in groups I and III. Based on the data, it can be stated that low plasma testosterone levels are related to higher collagen fiber density and active 3-caspase percentages in the bladder walls of orchiectomized and senile rats, respectively.

References

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Specify source of funding or grant	None
Is this a clinical trial?	No
What were the subjects in the study?	ANIMAL
Were guidelines for care and use of laboratory animals followed	Yes
or ethical committee approval obtained?	
Name of ethics committee	Federal University of Sao Paulo Animal Ethics Committee