

Journal of Andrology, Vol. 26, No. 6, November/December 2005  
Copyright © [American Society of Andrology](#)  
DOI: 10.2164/jandrol.04156

# Tungstate Treatment Improves Leydig Cell Function in Streptozotocin-Diabetic Rats

JOAN BALLESTER<sup>\*</sup>, JORGE DOMÍNGUEZ<sup>†</sup>, M. CARMEN MUÑOZ<sup>†</sup>,  
MERITXELL SENSAT<sup>\*</sup>, TERESA RIGAU<sup>\*</sup>, JOAN J. GUINOVAR<sup>†</sup> AND  
JOAN E. RODRÍGUEZ-GIL<sup>\*</sup>

*From the <sup>\*</sup> Unit of Animal Reproduction, Department of Animal Medicine and Surgery, School of Veterinary Medicine, Autonomous University of Barcelona, Bellaterra, Spain; and the <sup>†</sup> Department of Biochemistry and Molecular Biology and IRBB, Barcelona Science Park University of Barcelona, Barcelona, Spain.*

Correspondence to: Dr Joan E. Rodríguez-Gil, Unit of Animal Reproduction, Department of Animal Medicine and Surgery, School of Veterinary Medicine, Autonomous University of Barcelona, E-08193 Bellaterra, Spain.

Oral administration of sodium tungstate to adult male streptozotocin-diabetic rats for 3 months normalized serum levels of glucose, insulin, luteinizing hormone, and follicle-stimulating hormone. These effects were accompanied by an increase in reproductive performance, which was related to a strong improvement in Leydig cell function markers, such as the recovery of the number of Leydig cells and serum testosterone levels. Moreover, this in vivo recovery was related to a concomitant increase in the cell expression of insulin receptors. Tungstate treatment did not modify Leydig cell function in healthy rats. Furthermore, the addition of tungstate or insulin to the mTLC-1 cell line from Leydig cell origin increased the phosphorylation states of MAP-kinase and glycogen synthase kinase-3. Our results indicate that tungstate treatment in diabetic rats leads to a recovery of reproductive performance by increasing the number of Leydig cells. This increase contributes to the recovery of their functionality, thereby improving the overall function of these cells. We propose that this improvement is caused by the combined effect of the tungstate-induced normalization of insulin glucose and luteinizing hormone serum levels and a direct action of the effector on Leydig cells through modulation of at least MAP-kinase and glycogen synthase kinase-3 activities.

Key words: Diabetes, male, reproductive performance

This article has been cited by other articles:

## This Article

- ▶ [Full Text](#)
- ▶ [Full Text \(PDF\)](#)
- ▶ [Alert me when this article is cited](#)
- ▶ [Alert me if a correction is posted](#)

## Services

- ▶ [Similar articles in this journal](#)
- ▶ [Similar articles in PubMed](#)
- ▶ [Alert me to new issues of the journal](#)
- ▶ [Download to citation manager](#)

## Citing Articles

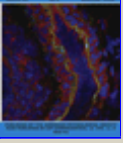
- ▶ [Citing Articles via HighWire](#)
- ▶ [Citing Articles via Google Scholar](#)

## Google Scholar

- ▶ [Articles by Ballester, J.](#)
- ▶ [Articles by Rodríguez-Gil, J. E.](#)
- ▶ [Search for Related Content](#)

## PubMed

- ▶ [PubMed Citation](#)
- ▶ [Articles by Ballester, J.](#)
- ▶ [Articles by Rodríguez-Gil, J. E.](#)



M. Miro-Queralt, J. J. Guinovart, and J. M. Planas  
Sodium tungstate decreases sucrose and Na<sup>+</sup>/D-glucose  
cotransporter in the jejunum of diabetic rats

Am J Physiol Gastrointest Liver Physiol, September 1, 2008; 295(3): G479  
- G484.

[\[Abstract\]](#) [\[Full Text\]](#) [\[PDF\]](#)