

## · 临床论著 ·

血清  $\beta$ -半乳糖凝集素-3 在非创伤股骨头坏死中的意义<sup>△</sup>陈敏<sup>1</sup>, 刘珊<sup>2</sup>, 王荣<sup>2</sup>, 刘歆<sup>3\*</sup>

(1. 青岛市胸科医院检验科, 山东青岛 266043; 2. 青岛大学附属青岛市中心医院, 青岛肿瘤医院检验科, 山东青岛 266042; 3. 临沂市人民医院股骨头科, 山东临沂 276000)

**摘要:** [目的] 本研究旨在探讨血清  $\beta$ -半乳糖凝集素-3 (lectin galactoside-binding soluble 3, LGALS3) 水平在非创伤性股骨头坏死 (non-traumatic osteonecrosis of the femoral head, NONFH) 诊断中的临床价值。[方法] 2023 年 4 月—2023 年 9 月纳入 NONFH 患者 84 例为坏死组, 选取同期体检者健康人 78 例为对照组。采用酶联免疫吸附法检测两组血清中 LGALS3 的水平, 并收集相关临床资料, 包括性别、年龄、体重指数 (Body Mass Index, BMI)、吸烟史、NONFH 病因、受累侧数、疼痛视觉模拟评分 (Visual Analogue Scale, VAS)、Harris 评分、坏死程度及 ARCO 分期。对两组 LGALS3 浓度及其与各临床参数的关系进行比较分析。[结果] 坏死组血清中 LGALS3 浓度显著高于正常人组 [(9.8±7.7) ng/ml vs (4.2±4.1) ng/ml,  $P<0.01$ ]。在 NONFH 患者中, 双侧受累组的 LGALS3 浓度显著高于单侧受累组 [(14.7±8.) ng/ml vs (5.3±3.3) ng/ml,  $P<0.001$ ], 股骨头塌陷者显著高于未塌陷者 [(13.5±7.3) ng/ml vs (3.1±1.2) ng/ml,  $P<0.001$ ]。根据 ARCO 分期, 从 I 期至 IV 期 LGALS3 浓度呈逐渐升高趋势, 且各亚组间差异有统计学意义 [(2.1±1.0) ng/ml vs (4.1±1.3) ng/ml vs (11.0±5.8) ng/ml vs (16.8±8.2) ng/ml,  $P<0.001$ ]。NONFH 患者的 LGALS3 浓度与 VAS 评分呈显著正相关 ( $r=0.843$ ,  $P<0.001$ ), 与 Harris 评分呈显著负相关 ( $r=-0.710$ ,  $P<0.001$ ), 与 ARCO 分期呈显著正相关 ( $r=0.822$ ,  $P<0.001$ )。ROC 分析表明, LGALS3 浓度预测是否 NONFH 的曲线下面积 (area under curve, AUC) 为 0.769。[结论] 血清中的 LGALS3 水平可反映 NONFH 病情的严重程度, 并可能作为该病早期诊断的潜在生物标志物。

**关键词:** 非创伤性股骨头坏死,  $\beta$ -半乳糖凝集素-3, 酶联免疫吸附法**中图分类号:** R681.18 **文献标志码:** A **文章编号:** 1005-8478 (2024)

**Significance of serum  $\beta$ -galectin-3 in non-traumatic necrosis of the femoral head** // CHEN Min<sup>1</sup>, LIU Shan<sup>2</sup>, WANG Rong<sup>2</sup>, LIU Xin<sup>3</sup>. 1. Department of Laboratory Medicine, Qingdao Chest Hospital, Qingdao 266043, China; 2. Department of Laboratory Medicine, Qingdao Central Hospital, Qingdao University, Qingdao 266042, China; 3. Department of Femoral Head, People's Hospital of Linyi City, Linyi 276000, China

**Abstract:** [Objective] To investigate the clinical value of serum  $\beta$ -galactoside-binding soluble 3 (LGALS3) in the diagnosis of non-traumatic osteonecrosis of the femoral head (NONFH). [Methods] From April 2023 to September 2023, 84 NONFH patients were enrolled as necrosis group, while other 78 healthy subjects were selected as control group. The serum LGALS3 concentration in the two groups were detected by enzyme-linked immunosorbent assay, and related clinical data were collected, including gender, age, body mass index (BMI), smoking history, NONFH etiology, number of affected sides, visual analogue scale (VAS) for pain, Harris score, and ARCO staging. The LGALS3 concentration and its relationship with clinical parameters in the groups divided by different factors were compared and analyzed. [Results] The serum LGALS3 concentration in the necrotic group was significantly higher than that in healthy group [(9.8±7.7) ng/ml vs (4.2±4.1) ng/ml,  $P=0.01$ ]. In patients with NONFH, LGALS3 concentration was significantly higher in the bilateral affected than in the unilateral involved [(14.7±8.) ng/ml vs (5.3±3.3) ng/ml,  $P<0.001$ ], the femoral head collapsed was significantly higher than the non-collapsed [(13.5±7.3) ng/ml vs (3.1±1.2) ng/ml,  $P<0.001$ ]. Based on ARCO staging, the concentration of LGALS3 gradually increased from stage I to stage IV, with statistically significant differences among the subgroups [(2.1±1.0) ng/ml vs (4.1±1.3) ng/ml vs (11.0±5.8) ng/ml vs (16.8±8.2) ng/ml,  $P<0.001$ ]. LGALS3 concentration in NONFH patients was significantly positively correlated with VAS score ( $r=0.843$ ,  $P<0.001$ ), whereas significantly negatively correlated with Harris score ( $r=-0.710$ ,  $P<0.001$ ), and positively correlated with ARCO stage ( $r=0.822$ ,  $P<0.001$ ). ROC analysis showed that the area under curve (AUC) of LGALS3 concentration predicting NONFH was 0.769. [Conclusion]

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作者简介: 陈敏, 主管技师, 研究方向: 疑难疾病的诊断性检验, (电话) 13969740233, (电子信箱) 13969740233@139.com

\* 通信作者: 刘歆, (电话) 15168917726, (电子信箱) linxin\_3610@126.com