Elevated SOX-9 Expression as a Biomarker for Lung Cancer Progress and Chemotherapy

for Lung Cancer Progress and Chemotherapy
Response: A Combined Analysis of PBMC

and Tissue Levels

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Introduction

Lung cancer, a leading global cause of cancer mortality, urgently requires non-invasive biomarkers to monitor progression and predict chemotherapy resistance(1). SOX9, a transcriptional driver of metastasis and chemoresistance(2), has been studied only in tumor tissue, neglecting its potential in peripheral blood. This study pioneers dual SOX9 analysis in PBMCs and tumor tissue, bridging this gap to establish its role as a novel biomarker for real-time monitoring and personalized therapy

Material and Methods

 30 lung cancer patients and matched healthy controls explored for SOX9 as a dual tissue/blood biomarker.

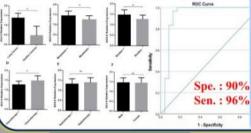


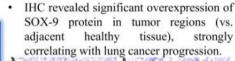


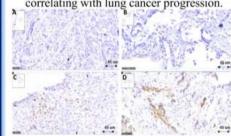


Results

SOX-9 expression in PBMCs was significantly upregulated in lung cancer patients compared to healthy controls, correlating with disease progression and chemotherapy response.







Discussion

RhoA GTP FAK Src Can channel Adenylate cyclase RN 1738 F-actin ROCK PAK Src Can channel Adenylate cyclase RN 1738 ROCK PAK Src Can channel Adenylate cyclase RN 1738 ROCK PAK Src Can channel Adenylate cyclase RN 1738 ROCK PAK Src Can channel ROCK PAK ATP PRA ATP PR

Conclusion

Combined analysis of SOX-9 in PBMCs and lung tissue identifies it as a precise biomarker for predicting cancer progression, metastasis, and therapy response, enabling personalized treatment strategies.

References

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