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EUROPEAN SOCIETY OF HUMAN GENETICS

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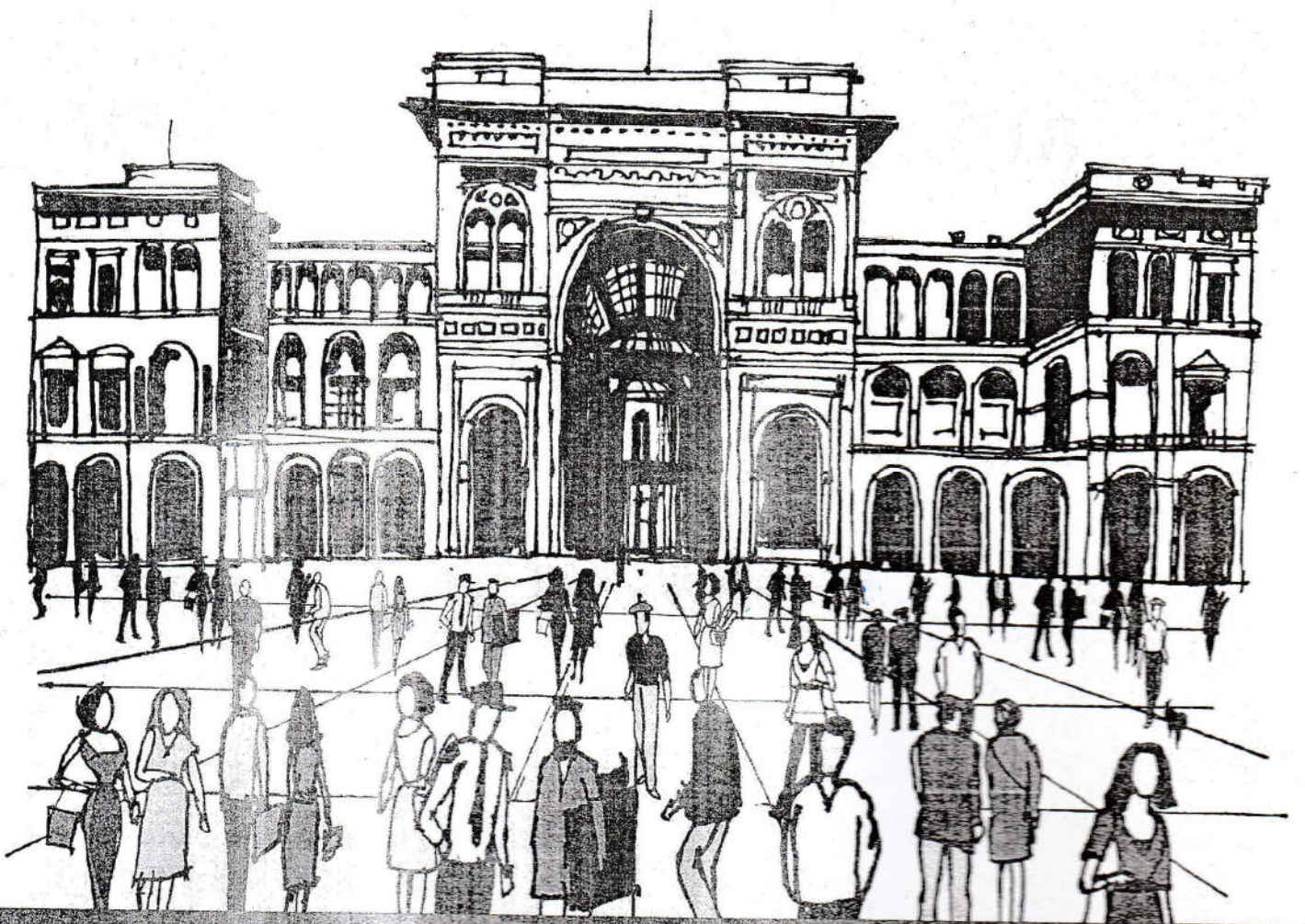
EUROPEAN MEETING ON PSYCHOSOCIAL ASPECTS OF GENETICS

**THE EUROPEAN SOCIETY
OF HUMAN GENETICS**

EUROPEAN HUMAN GENETICS CONFERENCE 2018

in conjunction with the European Meeting
on Psychosocial Aspects of Genetics

MiCo | Milan - Italy | June 16 - 19



PROGRAMME

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Session P12 - Cancer genetics



Itinerary

P12.176D / D - Diagnostic competence of mir-375 and mir-93-5p serum levels in malignant and non-malignant prostate diseases

📅 June 18, 2018, 4:45 PM - 5:45 PM

📍 Poster Area

Authors

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Disclosures

Y. Dulgeroglu: None. O. Eroglu: None.

Abstract

Introduction: The study purpose is to examine the diagnostic competence of miR-375 and miR-93-5p serum levels in prostate cancer differential diagnosis. It is also aimed to examine whether there is an interfering situation in the differentiation of chronic prostatitis, benign prostatic hyperplasia (BPH) and prostate cancer, considering the oncogenic and tumor suppressor properties of miRNAs together with the proinflammatory characteristics.

Materials and Methods: 25 Patients with BPH, 10 patients with chronic prostatitis and 33 patients with prostate cancer were included in the study. RNA isolation, cDNA synthesis and qRT-PCR steps were performed with Qiagen branded kits based on SYBR Green method, using the protocol of the commercial company. For statistical analysis, $-\Delta\text{Ct}$ values, obtained from the use of ce-miR-39 Ct values in normalization, were used. Differences between groups were tested by independent sample t test and variance analysis. ROC analysis was performed to calculate the diagnostic competence. "Fold change" calculations were performed online on Qiagen webpage. In all analyzes, alpha error level was accepted as 0.05. **Results:** The results are summarized in Table-1. **Conclusion:** It was observed that miR-375 and miR-93-5p differ in significantly between malignant and non-malignant disease groups. It was assessed that the sensitivity of miR-375 to differentiate non-malignant diseases from malignant diseases was higher than prostate specific antigen (PSA) and chronic prostatitis may be an interfering condition in BPH and cancer differential diagnosis.

Results

Compared Groups	miRNA	P value	Fold Change	AUC	Specificity	Sensitivity
Malignant-Non-malignant	miR-375	<0.001	-4.48	0.781	91%	46%
Malignant-Non-malignant	miR-93-5p	0.045	-1.79	0.662	91%	23%
BPH-Chronic prostatitis	miR-375	0.324	-	-	-	-
BPH-Chronic prostatitis	miR-93-5p	0.338	-	-	-	-
BPH-Cancer	miR-375	<0.001	-5.60	0.829	91%	56%
BPH-Cancer	miR-93-5p	0.392	-	-	-	-
Chronic prostatitis-cancer	miR-375	0.169	-	-	-	-
Chronic prostatitis-cancer	miR-93-5p	0.046	-2.78	0.744	91%	50%