Abstract 3029: Plasma hPG₈₀ (circulating Progastrin) levels in cancer patients in Nigeria: Prolevcan study

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Background:

Progastrin is a tumor-promoting peptide which is detectable in the blood of patients with different cancers. hPG_{80} (circulating progastrin) is produced by cancer cells. Recently, it was reported that hPG_{80} is detected in the blood of cancer patients, suggesting its potential utility for cancer detection. In this Nigerian study, we assessed the performance of hPG_{80} in diagnosed cancer patients versus healthy volunteers.

Methods:

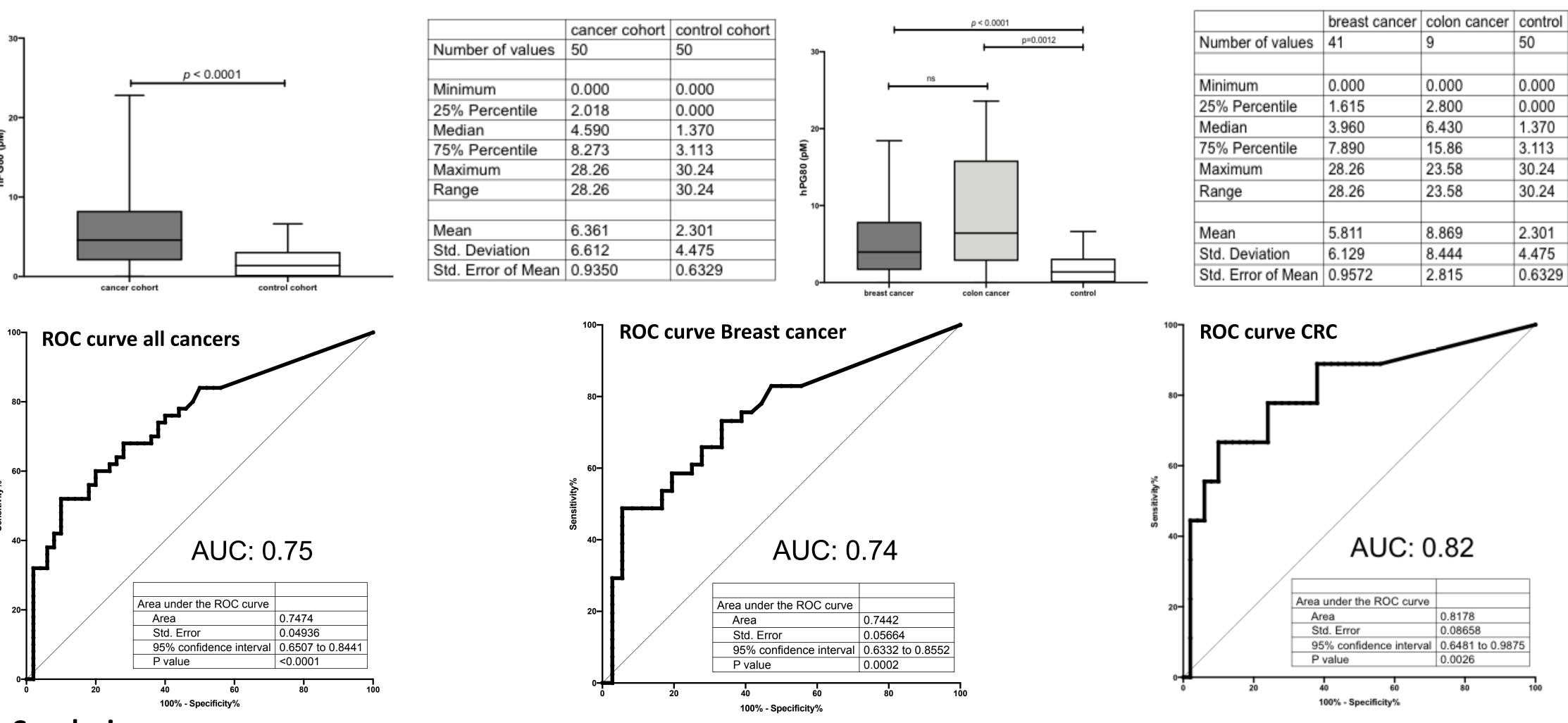
Plasma samples of 50 patients with breast (n=41) and colorectal (n=9) cancer, aged from 26 to 70 years, were assayed for hPG₈₀ levels with the DxPG₈₀ kit from ECS-Progastrin. The diagnostic performance (ROC AUC) of hPG₈₀ was assessed compared to 50 healthy volunteers aged from 21 to 38 years.

		Breast cancer N n = 41	CRC N n = 9	N n = 50
Age, years	Median (range)	50 (27-70)	57 (26-70)	29 (18-38)
Gender	Male	0	4	14
	Female	41	5	36
Menopause		19	/	0
Histological type	Ductal carcinoma	34	A 543 9	
	Others	8		
Immunohistochemical profile	Triple negative	11		
	HR positive	10	/	
	Other or unknown	20		
Clinical stage	1	2	8	/
	11	4		
	111	35	5	
	IV	0	3	

Clinical and pathological characteristics

Results:

Plasma hPG₈₀ levels were significantly higher in cancer patients compared to controls (median values: 4.59 pM (IQR: 2.02-8.27 pM) vs 1.37 pM (IQR: 0-3.11 pM), p < 0.0001). The median value of hPG₈₀ level was 3.96 pM (IQR: 1.61-7.89 pM) for breast cancers and 6.43 pM (IQR: 2.80-15.86 pM) for colorectal cancer (CRC) patients. ROC AUC for all cancers, breast cancer and colorectal cancer were 0.75, 0.74 and 0.82, respectively. There was no correlation between hPG₈₀ blood levels and age or CA15.3 levels.



All cancers combined cohort vs control

Conclusions:

Plasma hPG₈₀ is a simple and relatively affordable blood test, it shows potential utility as a biomarker for cancer detection, monitoring and treatment assessment.

Further prospective studies are needed to explore and confirm its potential.

Breast and CRC cancer cohorts vs control