Correction to: Impact of new generation hormone-therapy on cognitive function in elderly patients treated for a metastatic prostate cancer: Cog-Pro trial protocol

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Correction

After publication of the original article [1] the authors found that Table 2 had been formatted incorrectly, meaning that some rows in the Table did not display the correct information.

An updated version of Table 2 is included with this Correction.

The original article has also been updated.

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Table 2 Used cognitive tests, questionnaires and biological tests

Evaluations	Before inclusion	At inclusion (baseline) [b]	3 months	6 months	12 months
Signed Informed Consent	✓				
Previous medical history	✓				
Cognitive assessment [a] MoCA Grober-Buschke test Digit span forward and backward (WAIS-III) Code (WAIS III) Trail Making test Doors test Stroop Victoria Verbal fluencies Rey-Osterrieth Complex Figure Number location (VOSP) Years of education and fNART		✓ ✓ Only at inclusion	✓	✓	✓
Quality of life FACT-G, FACIT-F, FACT-Cog, HADS, ISI		✓	✓	✓	✓
Pain (VAS)	✓ [c]	✓	✓	✓	✓
ONLY for PATIENTS (group of interest and control group)					
Geriatric assessment [d] G8 Charlson ADL IADL MNA Time up and go		✓	✓	✓	✓
Quality of life FACT-P		✓	✓	✓	✓
Adherence evaluation [e] Morisky questionnaire Patient diary			✓	✓	✓
Biological tests [f]		✓	✓	✓	✓
Specific blood samples for further research [g]		✓			

MoCA Montreal Cognitive Assessment, *WAIS* Wechsler Adult Intelligence Scale, *VOSP* Visual Object and Space Perception Battery, *fNART* French National Adult Reading Test, *ISI* Insomnia Severity Index, *VAS* Visual Analog Scale, *ADL* Activities of Daily Living, *IADL* Instrumental Activities of Daily Living, *MNA* Mini-Nutritional Assessment

[a]Cognitive assessment will be performed by neuropsychologists

[b]For group of interest patients: before the start of the treatment or within 15 days after the start of treatment by abiraterone acetate or enzalutamide

[c]Had to be ≤3 on the 0–10 pain VAS scale to meet with inclusion pain criteria

[d]Geriatric assessment will be performed by a study nurse specialized in geriatric

[e]Adherence evaluation will be performed only in group of interest patients

[f]At each time: CBC, platelets, albumin, CRP, prealbumin, iron, ferritin, transferrin, creatinin, sodium, potassium, ALT, AST, GGT, ALP, total bilirubin, TSH, T4, testosterone. At inclusion only: cortisol (at 8 h AM, fasting)

[g]1 EDTA (5 ml), 1 dry tube with gel (5 ml) and 1 dry tube without gel (5 ml)