(15.3%), psychotic disorders (4%) and drug use (2%). There were no patients with eating or conduct disorders or IPI.

**Conclusions**

Psychiatric morbidity is frequent in resistant-epilepsy. Despite 38% of patients suffered from at least one axis I diagnoses, IDD was the most prevalent condition and not included in SCID interview.

**Disclosure of interest**

The authors have not supplied their declaration of competing interest.

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**EW0620**

**Cannabinoid hyperemesis syndrome, a treatment discussion**


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**Introduction**

Cannabinoid hyperemesis syndrome (CHS), is characterized by recurrent episodes of severe nausea and intractable vomiting, preceded by chronic use of cannabis. A pathognomonic characteristic is compulsive bathing in hot water. The resolution of the problem occurs when cannabis use is stopped. However, patients are often reluctant to discontinue cannabis. Treatment with anti-emetic medication is ineffective. Case series suggested haloperidol as a potential treatment. Other antipsychotics as olanzapine has been used as anti-emetic treatment in chemotherapy.

**Objectives**

To describe three cases of patients with CHS whom showed a successful response to olanzapine, even when, haloperidol had failed.

**Aims**

To present an alternative treatment for CHS which can offer benefits over haloperidol.

**Methods**

We present three cases of patients who suffered from CHS and were admitted to emergency department. All patients were treated with olanzapine after conventional anti-hemetic treatment failure. One patient was also unsuccessfully treated with haloperidol.

**Results**

All three patients showed a good response to olanzapine treatment. Different presentations were effective: velotub and intramuscular. Their nausea, vomits and agitation were alleviated. They could be discharge after maintained remission of symptoms.

**Conclusions**

Olanzapine should be considered as an adequate treatment for CHS. Its suitable receptorial profile, its availability in different routes of administration and its side effects profile could offer some benefits over haloperidol.

**Disclosure of interest**

The authors have not supplied their declaration of competing interest.

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**EW0621**

**An Italian observational study on subclinical cardiovascular risk factors and depressive symptomatology. A suggestion for the potential utility of a sinergic cardio-psychiatric perspective**


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**Aim**

Aim of this study was to investigate the possible link between subclinical CV risk factors (atherosclerosis), depressive symptoms, and inflammation.

**Methods**

Cross-sectional study. Inclusion criteria: outpatients aged ≥40 years, attending colonoscopy after positive faecal occult blood test, negative medical history for cancer. Collected data: blood pressure, glycaemia, lipid profile, waist circumference, BMI, PCR (C reactive protein), LPS (bacterial lipopolysaccharide), ultrasound carotid intima-media thickness (c-IMT), Psychometric tests: HADS, TCI, IMSA, SF36. Statistical analysis performed with STATA13.

**Results**

The 54 patients enrolled were equally distributed by gender. CV risk factors were common in the study population, with 33 patients (61.11%) with hypertension, 14 (25.93%) with hyperglycemia, 20 (37.4%) with hypertriglyceridemia, 19 (35.19%) with low HDL and 64.81% with overweight. High levels of PCR were found in 24 subjects (44.44%). Right c-IMT was increased in 26.41% of the sample, and 11.32% had an atheromatous plaque. Left c-IMT was increased in 24.53% of patients, with a plaque in 7.55% of them. Clinically relevant depressive symptoms were found in the 18.87% of the sample and were statistically significantly associated with PCR (OR = 28.63; P = 0.01).

**Conclusions**

Evidence contributing to the so-called “inflammation theory” of depression and supporting the association between mood and CV disorders was here collected, supporting the need for a multidisciplinary approach to the diagnosis and treatment of such conditions, assuming a clinically-translated PNEI (psycho-neuro-endocrino-immunological) perspective.

**Disclosure of interest**

The authors have not supplied their declaration of competing interest.

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