LIVERPOOL

The aim of this study was to determine the predictive and/or prognostic value of progastrin, in patients with HCC, before and after curative treatment (surgical resection, radiofrequency ablation) or palliative treatment (transcatheter arterial chemoembolization, targeted therapy, and immunotherapy)

METHODS:

This study is a prospective cohort monocentric trial in CHU Montpellier.

1- inclusion criteria:

- * Male or female ≥ 18 years old
- * Histologically confirmed diagnosis HCC
- * HCC suspected in RMI and/or scanner
- * In full capacity of their psychological condition, familial, sociological or potentially unfavorable geographical agreement to great study protocol and monitoring program.
- * Patient information and written informed consent form signed

2- excluded criteria:

- * Patient with an other cancers in last 5 years
- * Current infection
- * Patient with autoimmune disease (as lupus, IBD, scleroderma, ...
- * top athlete

3- Methods:

The assay is performed from a fasting blood sample in EDTA tube (2 test tubes of 7 mL per sample).

The samples will be transferred to Biochemical department for anonymization, centrifugation (2000g during 15 minutes) and freezing (- 80°C).

The biobank was declared since 2014 december at CHU Montpellier (n° BB-0033-00031).

4- Monitoring

The biobank openning after once the relevant ethics committee has given its authorisation (N° 2019_IRB-MTP_01-11- MR004)

Patient follow up is every 2 month until death.