

COVIPACT - Impact of the COVID-19 epidemic on patient management in the oncology-haematology setting, and on psychological repercussions among patients and caregivers

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Last update : 04/30/2021 | Version : 1 | ID : 73892

General

Identification

| | |
|-----------------|---|
| Detailed name | Impact of the COVID-19 epidemic on patient management in the oncology-haematology setting, and on psychological repercussions among patients and caregivers |
| Sign or acronym | COVIPACT |

General Aspects

| | |
|-----------------------------------|------------------------|
| Medical area | Cancer research |
| Study in connection with Covid-19 | Yes |
| Pathology (details) | CANCER |
| Health determinants | Lifestyle and behavior |

Scientific investigator(s) (Contact)

| | |
|----------------------|-----------------------------------|
| Name of the director | FAVEYRIAL |
| Surname | Audrey |
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| Organization | FRANCOIS BACLESSE CENTRE |

Collaborations

| | |
|---|----|
| Participation in projects, networks and consortia | No |
|---|----|

Funding

Governance of the database

Sponsor(s) or organisation(s) responsible FRANCOIS BACLESSE CENTRE

Organisation status Both

Presence of scientific or steering committees No

Additional contact

Name of the contact LECONTE

Surname Alexandra

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Email a.leconte@baclesse.unicancer.fr

Organization François Baclesse Centre

Main features

Type of database

Type of database Others

Specify data obtained from medical records

Study databases (details) Cohort study

Database recruitment is carried out by an intermediary A selection of health institutions and services

Database recruitment is made on the basis of: Another treatment or procedure

Database recruitment is carried out as part of an interventional study No

Additional information regarding sample selection. Recruitment of patients and healthcare professionals

Database objective

Main objective Evaluate the impact of the COVID-19 pandemic on adjustments to medical cancer treatment delivered in the outpatient setting to patients with cancer or haematological malignancies undergoing treatment.

| | |
|--------------------|---|
| Inclusion criteria | <ul style="list-style-type: none"> - Adult patient, treated for a solid tumour or haematological malignancy - Receiving or needing to receive medical cancer treatment delivered in an outpatient oncology setting at participating sites: treatment initiated before or during the COVID-19 pandemic - The patient does not object to taking part in the study - Patient not deprived of their liberty or under supervision - Female patient with no associated geographical, social or psychopathological condition liable to compromise the patient's ability to take part in the study |
|--------------------|---|

Population type

| | |
|-----|---|
| Age | <ul style="list-style-type: none"> Adulthood (19 to 24 years) Adulthood (25 to 44 years) Adulthood (45 to 64 years) Elderly (65 to 79 years) Great age (80 years and more) |
|-----|---|

| | |
|--------------------|-----------------|
| Population covered | Sick population |
|--------------------|-----------------|

| | |
|-----------|-------------------------------|
| Pathology | C00-C97 - Malignant neoplasms |
|-----------|-------------------------------|

| | |
|--------|---|
| Gender | <ul style="list-style-type: none"> Male Woman |
|--------|---|

| | |
|----------------|----------|
| Geography area | Regional |
|----------------|----------|

| | |
|--|-----------|
| French regions covered by the database | Normandie |
|--|-----------|

Data collection

Dates

| | |
|--|------|
| Date of first collection (YYYY or MM/YYYY) | 2020 |
|--|------|

| | |
|---|------|
| Date of last collection (YYYY or MM/YYYY) | 2021 |
|---|------|

Size of the database

| | |
|--|------------------------|
| Size of the database (number of individuals) | [500-1000[individuals |
|--|------------------------|

| | |
|--------------------------------------|-----|
| Details of the number of individuals | 808 |
|--------------------------------------|-----|

Data

Database activity Current data collection

Type of data collected Clinical data

Clinical data (detail) Direct physical measures

Presence of a biobank No

Procedures

Participant monitoring Yes

Monitoring procedures Monitoring by contact with the participant (mail, e-mail, telephone etc.)

Links to administrative sources No

Promotion and access

Promotion

Access