

Gilles DULUC

Directeur de la Recherche Clinique
et de l'Innovation

Talence, le 13/05/2025

Jeanne PATARD

Directrice adjointe de la Recherche
Clinique et de l'Innovation

Département Promotion Interne

Anne GIMBERT

Responsable
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Object : Interventional Clinical Trial's results posting to the European Clinical Trial register (EudraCT) of the European Medicines Agency (EMA).

In accordance of Directive 2001/20/CE

Study Title : MISOBOLD : Prostate cancer hypoxia using BOLD MRI and 18F-FMISO PET imaging

EudraCT Nbr : 2015-000377-12

Sponsor Nbr : CHUBX2014/26 / **ANSM Nbr :** 170339A-13

Transitioned study to the Regulation (EU) 536/2014 : no


Status of the study : Prematurely ended study

The study has been prematurely ended because of a staff turnover through the research staff. As a result the end of the regulatory authorization has been reached. No patient has been recruited in the study and the steering committee decided to close the study.

For the Head of the University Hospital of Bordeaux
and by delegation,
The Director of the Clinical research and Innovation

Gilles DULUC

par délégation Jeanne Patard, directrice adjointe



DIRECTION GENERALE

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