Institutional Conflict of Interest Management and Monitoring Plan: Ipsen Bioscience, Inc.

The University of Texas MD Anderson (MD Anderson) and Ipsen Bioscience, Inc. (Ipsen) are parties to a Patent and Technology License Agreement and Collaboration Agreement (Agreements) pursuant to which MD Anderson will conduct a first Phase I clinical trial and any other clinical trial for Licensed Technology that MD Anderson and Ipsen agree to include in the Research Plan, as well as development activities listed in the Research Plan (Studies).

Under the Agreements, MD Anderson has a right to funding from Ipsen for MD Anderson's conduct of the Studies. Such royalties and payments, including the royalty payments to be paid to MD Anderson by Ipsen upon the first dosing of the first patient on various clinical trials with Licensed Technology create an institutional conflict of interest.

Because MD Anderson is committed to the protection of human subjects and the effective management of its financial conflict of interest in relation to its research activities, MD Anderson has implemented an Institutional Conflict of Interest Management and Monitoring Plan (Plan) to manage and monitor the conflict of interest with respect to MD Anderson's conduct of the Studies. The Plan has been approved by the President of MD Anderson and the Executive Vice Chancellor for Health Affairs for The University of Texas System (EVC) and has been implemented by MD Anderson.

The Plan requirements include:

- MD Anderson employees, who have a financial interest in Ipsen and will be involved in the conduct of the Studies, will have a personal conflict of interest management plan covering their involvement of the Studies.
- Disclosure of MD Anderson's financial interest and the financial interest of individual creators and inventors of the Licensed Technology involved in the conduct of the Studies, including a disclosure of the percentage share of payments payable to such individuals, to participants in the Studies,
- Disclosure of MD Anderson's financial conflict of interest, the financial interest of any individual creators and inventors of the Licensed Technology involved in the conduct of the Studies, and the Plan to all members of the research teams who will work on the Studies,
- Disclosure of MD Anderson's financial conflict of interest in all publications and oral presentations concerning the Studies,
- Supply information about the consent process that will include observation by a non-MD Anderson external ethicist of the consent process for the first patients who will receive the first dosing on Clinical Trials following to the External IRB,
- Review and confirm the first patient's Clinical Trial eligibility by a non-MD Anderson oncologist with appropriate expertise and no financial interests in Ipsen, prior to administration of the first dosing of the first patients on Clinical Trials upon which MD Anderson will receive milestone payments from Ipsen,
- Posting of this summary on MD Anderson's public website,
- Referral of any concerns/complaints related to MD Anderson's compliance with the Plan, or its financial conflict of interest, to The University of Texas System,
- Recusal of any MD Anderson Institutional Decision Maker who has a financial relationship with Ipsen or its known affiliates from negotiations with respect to any agreements or purchasing decisions,
- Oversight of Studies by an external Institutional Review Board (External IRB), including reporting to the External IRB by MD Anderson's Investigational New Drug (IND) Office when applicable,
- Engagement of a non-MD Anderson ethicist (External Ethicist) to address any questions or concerns that participants in the Studies may have pertaining to the MD Anderson financial interest and conflict of interest,
- Supply a copy of the Plan to the External IRB and External Ethicist,
- Review of safety and efficacy data of Studies that are clinical trials by an external and independent Data Safety Monitoring Board (External DSMB),

- Use of multi-institutional trials with a non-MD Anderson lead principal investigator for Studies that are Phase III or Phase II clinical trials aimed at gaining FDA approval under a new drug or biological license application,
- Monitoring activities related to the manufacture of Investigational Agents, if required for the Studies, by MD Anderson's IND Office,
- Reporting to the EVC by an External Contract Research Organization on Studies that are IND-enabling preclinical studies,
- Review and revision of the Plan as necessary with any amendments requiring EVC approval, and
- Annual review of MD Anderson's compliance with the Plan by The University of Texas Systemwide Compliance Officer, with a written report of the review from the External DSMB to be provided to the EVC.

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