### Please complete the ABSTRACT TEMPLATE online, for Biomed 2023 Company Presentations

# ABSTRACT TEMPLATE: CHECKLIST AND INSTRUCTIONS

All items marked with an \* are mandatory to complete The maximum number of words for this abstract is 400

### Please be sure to complete the following:

Company name Gilboa Therapeutics LTD \* Website www.gilboa.bio \*

CEO name Mr. Barry Labinger \* Cell phone number +1-301-768-8260 \*

Select a CATEGORY: Biotech/Pharma or Medical Devices or Health IT/Digital \* (Delete categories you are not selecting)

Select up to two SESSIONS per abstract from the list below \* (Delete sessions you are not selecting)

## 3. Are Cancer Therapeutics Fulfilling the Promise?

You may delete the section instructions, leaving **only the bolded bullet title Answers below should not exceed 60 words per question:** 

## • Executive Summary / Investment Rational

Gilboa Tx is a pre-clinical stage biotech company developing a breakthrough T cell therapy for solid cancers, that overcomes the inherent limitations of CAR T cells. These novel engineered T-cells, called **SolidT**, show effective and specific cytotoxicity only towards tumor cells, sparing normal tissues expressing low antigen levels, thus providing a favorable safety profile.

## • Core Technology

Peripheral blood T cells engineered with Gilboa's novel receptor, based on the highaffinity  $Fc\gamma RI$  scaffold, allowing T-cells to target solid tumor cells using antibody intermediates. This breakthrough technology manifests remarkable killing abilities against solid tumors in mouse models with an unprecedented safety profile.

# • Product Profile/Pipeline

Gilboa Tx is developing an extensive pipeline, including off-the-shelf **SolidT** based on engineered  $\gamma/\delta$  T cells, next-generation **SolidT** that target HER2 +1 tumors, and developing new combinations with proprietary antibodies to target diseases with urgent unmet medical need.

## • Business Strategy

The SolidT technology is an autologous cell therapy that will be developed as a therapeutic product in combination with relevant mAbs.

## • What's Next?

Round A funding, CMC activities and IND-enabling studies for our lead program toward Phase I clinical trial.