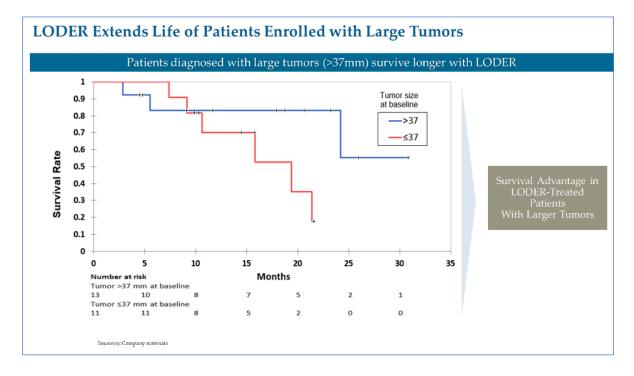
Dear Silenseed Shareholders,

With good Efficacy and Safety clinical results from Phase 2, Silenseed received FDA approval to initiate Phase 3, and are planning Kick-off at Q3-2021.

Clinical Development

Good news:

- KRAS-LODER converts more patients from non-Resectable to Resectable, 3 times better compared to control (27%. vs 9%).
- KRAS-LODER specifically affects patients with large tumors, showing a clear signal of LODER effect (figure).
- We continue to see very encouraging evidence of life extension, tumor response and pain reduction (Statistically significant reduction in pain was noted in the subgroup who enrolled with high pain score (VAS > 4)



With such a strong signal, both in safety and in efficacy, we received a green light from the FDA to initiate a larger (316 patients) Phase 3trial that aims to support marketing approval.

Silenseed

Quality:

We are scheduled to have a full quality inspection by the Israeli Minister of Health at April 25, 2021; that will support Phase 3 and later commercialization of our products (planned from 2024)

Team:

Silenseed has strengthened the team with great people including:

Dr Eran Gefen (TEVA) as our VP BD; Dr. Diana Gurevitch (VP Pre-clinical); Eran Ovadya (CFO), Patricia Erb (Clinical), Michal Yaron (VP Regulatory Affair)

I wish to thank you very much for your continuous support in Silenseed. Without your support nothing of the above could become a reality.

Sincerely,

Amotz Shemi Chief Executive Officer Silenseed LTD

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