



Living Cell Technologies Limited Company Announcement

LCT receives approval to expand DIABECCELL[®] NZ Trial

- NZ Minister of Health approves the addition of four patients to Phase II trial
- New patients add further rigour to trial by generating valuable additional data

4 August 2010: Sydney, Australia & Auckland, New Zealand. Living Cell Technologies Limited (ASX: LCT; OTCQX: LVCLY), a global company pioneering the development of a cell implant to treat diabetes, today reported that the Minister of Health has approved an addition to the New Zealand Phase II clinical trial of DIABECCELL[®] to include more patients. To date, eight insulin dependent diabetes patients have received the implants DIABECCELL[®], LCT's encapsulated porcine islets, and now another four patients will be enrolled into the dose ranging trial. The expansion of the New Zealand study has also been approved by the Regional Ethics Committee and the Data Safety and Monitoring Board. The Minister stated that he is satisfied that amending the original approval to include more patients will add further rigour to the study and may provide valuable additional information.

LCT has recently reported that in the current trial of eight New Zealand patients with unstable diabetes, all have shown the benefit of reduction or elimination of episodes of low blood glucose levels that are often life-threatening. The dramatic results to date showing DIABECCELL[®]'s ability to ameliorate this serious complication of diabetes, known as hypoglycaemic unawareness, are one key indicator of potential benefit to patients.

The first four patients received one implant of DIABECCELL[®] at the dose of 10,000 islet equivalents per kilogram body weight (IEQ/kg) without any reported adverse events attributable to the treatment. A second group of four patients has received a higher dose of 15,000 IEQ/kg, also with no significant adverse events attributed to the treatment. At this time, the follow up period is too short to assess efficacy with this second group.

The safety profile of DIABECCELL[®] continues to be confirmed and therapeutic benefit remains promising as the Company progresses with its dose ranging trials, aimed at identifying the most efficient dose to achieve optimal efficacy.

Prof Bob Elliott, LCT Medical Director said: "This approval allows us to administer up to 20,000 IEQ/kg in four patients which I expect is likely to be the maximum consideration for a single dosing. It will enable a future assessment of a possible commonly seen plateau effect in dosing which will allow us to understand dosing efficiencies for various patient indications. "

Dr Paul Tan, NZ Chief Executive Officer LCT added: "The Minister's approval to expand the New Zealand trial is part of LCT's clinical and commercial development plan to see DIABECCELL[®] advance to market. This additional patient information will be important when analysing the data to assess the necessary doses to meet various endpoints relevant to patients with unstable Type 1 diabetes."

DIABECCELL[®] is LCT's treatment designed to normalise the lives of people with insulin dependent diabetes. DIABECCELL[®] comprises encapsulated porcine insulin-producing cells (islets) that are implanted into the abdomen of patients using a simple laparoscopic procedure, and work by self-regulating and efficiently secreting insulin in the patient's body. LCT's breakthrough proprietary encapsulation technology, IMMUPEL, means that patients receiving DIABECCELL[®] treatment do not require immunosuppression after implantation.

- Ends -

For further information: www.lctglobal.com

<p>At the company: Dr Paul Tan Chief Executive Officer Tel: +64 9 276 2690 ptan@lctglobal.com</p> <p>Prof Bob Elliott Medical Director Tel: +64 9 276 2690 belliott@lctglobal.com</p> <p>Ms Susanne Clay Chief Business Officer Tel: +64 9 270 7954 sclay@lctglobal.com</p>	<p>Media and investor enquiries: NZ and Australia: Buchan Consulting Rebecca Wilson Tel: +61 3 9866 4722 Mob: +61 417 382 391 rwilson@bcg.com.au</p> <p>Paul Dekkers Tel: +61 2 9237 2800 pdekkers@bcg.com.au</p>
---	---

About Living Cell Technologies - www.lctglobal.com

Living Cell Technologies (LCT) is developing cell-based products to treat life threatening human diseases. The Company owns a biocertified pig herd that it uses as a source of cells for treating diabetes and neurological disorders. For patients with Type 1 diabetes, the Company transplants microencapsulated islet cells so that near-normal blood glucose levels may be achieved without the need for administration of insulin or at significantly reduced levels. The Company entered clinical trials for its diabetes product in 2007. For the treatment of Parkinson's disease and other neurological disorders, the company transplants microencapsulated choroid plexus cells that deliver beneficial proteins and neurotrophic factors to the brain. LCT's breakthrough encapsulation delivery technology, IMMUPEL, enables healthy living cells to be injected into patients to replace or repair damaged tissue without requiring the use of immunosuppressive drugs to prevent rejection. LCT also offers medical-grade porcine-derived products for the repair and replacement of damaged tissues, as well as for research and other purposes.

LCT Disclaimer

This document contains certain forward-looking statements, relating to LCT's business, which can be identified by the use of forward-looking terminology such as "promising," "plans," "anticipated," "will", "project", "believe", "forecast",

“expected”, “estimated”, “targeting”, “aiming”, “set to,” “potential,” “seeking to,” “goal,” “could provide,” “intends,” “is being developed,” “could be,” “on track,” or similar expressions, or by express or implied discussions regarding potential filings or marketing approvals, or potential future sales of product candidates. Such forward-looking statements involve known and unknown risks, uncertainties and other factors that may cause actual results to be materially different from any future results, performance or achievements expressed or implied by such statements. There can be no assurance that any existing or future regulatory filings will satisfy the FDA’s and other health authorities’ requirements regarding any one or more product candidates nor can there be any assurance that such product candidates will be approved by any health authorities for sale in any market or that they will reach any particular level of sales. In particular, management’s expectations regarding the approval and commercialization of the product candidates could be affected by, among other things, unexpected clinical trial results, including additional analysis of existing clinical data, and new clinical data; unexpected regulatory actions or delays, or government regulation generally; our ability to obtain or maintain patent or other proprietary intellectual property protection; competition in general; government, industry, and general public pricing pressures; and additional factors that involve significant risks and uncertainties about our products, product candidates, financial results and business prospects. Should one or more of these risks or uncertainties materialize, or should underlying assumptions prove incorrect, actual results may vary materially from those described herein as anticipated, believed, estimated or expected. LCT is providing this information and does not assume any obligation to update any forward-looking statements contained in this document as a result of new information, future events or developments or otherwise.