



Living Cell Technologies Limited Company Announcement

LCT Approved to Further Expand DIABECCELL[®] NZ Trial

- NZ Minister of Health approves addition of 2 patients to be dosed at 5,000 islet equivalents per kilogram of body weight
- New patients bring total number of study participants in NZ phase II trial to 14, spread across four dose groups
- Fourth treatment arm at low 5,000 dose will help build the dose ranging data set needed to define target product profile for Phase III trials

20 December 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 announced that the New Zealand Minister of Health has approved the addition of two patients to the New Zealand Phase II clinical trial of DIABECCELL. These patients will be in addition to the 12 patients already approved to receive DIABECCELL implants, LCT's encapsulated porcine islets for the treatment of Type 1 diabetes.

To date, 11 of the 12 approved New Zealand patients with unstable insulin dependent diabetes have received this ground-breaking treatment, which has been shown to improve diabetes management and reduce or eliminate episodes of life-threatening low blood glucose levels. The dramatic results to date show DIABECCELL's ability to ameliorate this serious complication of diabetes, known as hypoglycaemic unawareness, is an important potential benefit to patients.

The expansion of the trial will allow LCT to build its dose-ranging data set of DIABECCELL, which is key to determining the dose regimen which provides optimal patient benefit. The first four patients received one implant of DIABECCELL at a dose of 10,000 islet equivalents per kilogram body weight (IEQ/kg). A second group of four patients has received a higher dose of 15,000 IEQ/kg. In the third group of two patients, a high dose of 20,000 IEQ/kg was administered. The fourth group of four patients will receive a dose of 5,000 IEQ/kg.

The expansion of the New Zealand study has also been approved by the Northern X Regional Ethics Committee and the Data Safety and Monitoring Board. The safety profile of DIABECCELL continues to be confirmed and therapeutic benefit remains promising as the Company progresses with its dose-ranging studies and prepares for initiating pivotal Phase III trials next year.

Professor Bob Elliott, LCT Chairman and Medical Director said: "As often happens in medicine, we see a threshold zenith where there appears to be little additional therapeutic response to higher doses of DIABECCELL, and it is currently thought that this threshold may have been reached in this clinical trial. We believe reducing the dose may increase the clinical benefit of this treatment and will also mean we will require fewer islet cells to treat each patient. If this is proven true, it will be a very good outcome for the commercialisation of DIABECCELL."

Dr Ross Macdonald, Chief Executive Officer of LCT added: "The data generated from this trial is invaluable to finalizing our target product profile so we can finalise planning of our final Phase III trials next year. We are working quickly toward the commercialisation of DIABECCELL and gathering as much data as possible to ensure DIABECCELL offers the optimal level of benefit to patients."

- Ends -

For further information: www.lctglobal.com

At the company: Ms Susanne Clay Chief Business Officer, Living Cell Technologies Tel: +64 9 270 7954 Mobile: +64 21 418 833 sclay@lctglobal.com	Media and investor enquiries: NZ and Australia: Buchan Consulting Rebecca Wilson Tel: +61 3 9866 4722 Mobile: +61 417 382 391 rwilson@bcg.com.au Media Erik Denison Tel: +61 2 9237 2800 Mobile: +61 432 712 278 edenison@bcg.com.au
--	---

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 implants lead product DIABECCELL, 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, NTCELL, which delivers beneficial proteins and neurotrophic factors to the brain. LCT's breakthrough microencapsulation 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.