

Living Cell Technologies Limited Company Announcement

Appendix 4C Quarterly Cash Flow Report 31 December 2011

27 January 2012: Sydney, Australia & Auckland, New Zealand – Living Cell Technologies Limited (ASX: LCT; OTCOX: LVCLY), announced today the quarterly cash flow report for the quarter ended 31 December 2011. The Appendix 4C is attached. The cash balance at the end of the quarter was \$2,487,018 compared to \$2,875,447 at 30 September 2011. This decrease reflects the continuing development cost of the lead product DIABECELL[®] until 1 November and the primate study of NTCELL for the treatment of Parkinson's disease, partially offset by service fees.

Net operating cash outflows in the quarter were \$296,435 compared to \$1,797,312 last quarter. Receipts from customers were \$1,256,832 reflecting the services provided to Diatranz Otsuka Limited (DOL) from 1 November (last quarter \$3,589). Expenditure included \$984,148 on research and development (\$1,350,842 last quarter) for the supply of DIABECELL for the clinical trial in Argentina, services to DOL and preclinical studies of NTCELL. Cash received from grants was \$230,548 (\$230,924 last quarter).

Capital expenditure was \$0 in the quarter, compared to \$40,622 last quarter.

On 1 November 2011 the company settled the formation of a 50/50 company Diatranz Otsuka Limited (DOL) with Otsuka Pharmaceutical Factory, Inc. (OPF), to accelerate the commercialisation of DIABECELL. LCT transferred DIABECELL assets into Diatranz Otsuka Limited for \$25m of shares and OPF invested \$25m of cash in DOL. LCT provides research and development and administrative services to DOL at commercial rates and retains access to the facilities and designated pathogen free pigs for products other than diabetes.

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About Living Cell Technologies - <u>www.lctglobal.com</u>

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. The Company's lead product, DIABECELL[®], consists of microencapsulated porcine islets which are implanted into a patient's abdomen using a simple laparoscopic procedure. DIABECELL is designed to help normalise the lives of people with unstable Type 1 diabetes, especially those suffering from life-threatening episodes of unaware hypoglycaemia (low blood sugar), a dangerous and potentially fatal diabetes complication. The Company entered

clinical trials for its diabetes product in 2007 and very encouraging results have been reported to date. There have been no reports of remarkable product related adverse events. For the treatment of Parkinson's disease and other neurological disorders, the company implants microencapsulated choroid plexus cells NTCELL that deliver beneficial proteins and neurotrophic factors to the brain. LCT's technology 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.