FORM 51-102F3

MATERIAL CHANGE REPORT

Item 1 Name and Address of Company

PACIFIC THERAPEUTICS LTD. (the "Company")
Suite 1500 –409 Granville St. Vancouver, BC V6C 1T2
Telephone No. (604) 738-1049

Item 2 Date of Material Change

April 24, 2014

Item 3 News Release

A news release was disseminated April 24, 2014, posted to the CNSX website and was subsequently SEDAR filed with the securities commissions of British Columbia and Ontario.

Item 4 Summary of Material Change(s)

The Company announced that the Financial Industry Regulatory Authority (FINRA) has approved the Company's rule 15c2-11 submission as filed by the market maker Wilson-Davis & Co. The shares are quoted on the OTC Markets QB using the symbol PCFTF.

4.1 Full Description of Material Change

SEE PRESS RELEASE ATTACHED

4.2 Disclosure for Restructuring Transactions

Not applicable.

Item 5 Reliance on subsection 7.1(2) or (3) of National Instrument 51-102

Not applicable.

Item 6 Omitted Information.

Not applicable.

Item 7 Executive Officer

Doug Unwin,
President & CEO
604-738-1049

Item 8 Date of Report

April 24, 2014

FINRA CLEARS RULE 15C2-11 SUBMISSION AND ISSUES

SYMBOL – PCFTF

VANCOUVER, BC, Canada – April 24, 2014 – Pacific Therapeutics Ltd. (CSE: PT) (OTCQB: PCFTF) (Frankfurt: 1P3) (the "Company") is a clinical stage specialty pharmaceutical company focused on the repurposing and reformulation of existing FDA approved drugs for large markets. The Company's lead programs focus on erectile dysfunction and diseases of excessive scarring (fibrosis) which are $1 billion plus market opportunities.

The Company is pleased to announce that the Financial Industry Regulatory Authority (FINRA) has approved the Company's rule 15c2-11 submission as filed by the market maker Wilson-Davis & Co. The shares are quoted on the OTC Markets QB using the symbol PCFTF.

Doug Unwin CEO and President of the Company stated "having our shares quoted in the United States is the attainment of yet another of our stated milestones and is one step further towards our goal of increasing the Company's visibility in US equity markets as well as increasing the liquidity of the Company's shares".

ABOUT PACIFIC THERAPEUTICS LTD.

The Company's strategy includes reformulating approved drugs to increase efficacy and patient compliance, while reducing side effects, as well as completing the further clinical testing, manufacturing and other regulatory requirements sufficient to seek marketing authorizations. This strategy may reduce the risk, time and cost of developing therapies by avoiding the risks associated with basic research and using compounds with unknown safety and toxicity profiles.

Pacific Therapeutics Ltd. lead drug candidate for fibrosis (progressive scarring of the organ), PTL-202 is a combination of Pentoxifylline an FDA approved drug and N-Acetyl-Cysteine (NAC) an amino acid and an extremely potent and important antioxidant. The Company has completed an initial clinical trial of the combination with positive results. *"It's estimated that 45 % of all deaths are related to fibrotic changes" World Health Organization Feb, 2008.*

Worldwide, there are over 5,000,000 people living with Idiopathic Pulmonary Fibrosis (IPF), (IPF Coalition). IPF therapy sales across the US, France, Germany, Italy, Spain, and the UK to rise to over $1.1 billion by 2017, at a Compound Annual Growth Rate (CAGR) of 86.6% (RnR Market Research, 2013). IPF kills more patients per year than either prostate or breast cancer.

In 2011 the total market for drugs to treat erectile dysfunction ("ED") exceeded $5 billion. Pacific Therapeutics Ltd. has finalized a license to an oral dissolving technology ("sublingual formulation") of an approved drug to treat erectile dysfunction (ED).

Sales of the market leader alone exceeded $1.9 billion in 2011. The sublingual formulation may improve on existing drugs for erectile dysfunction potentially acting faster and with fewer side effects. As large pharmaceutical companies lose their patents on these drugs the opportunity has developed for innovative formulations of drugs for ED. This is a very exciting development for Pacific Therapeutics Ltd. as it shortens the time to market for the Company's first product and may add significantly to future revenues.

The Company plans to build on the already significant development of the sublingual treatment with the initiation of a pivotal Bioequivalence trial. With successful results from this trial the Company will begin the application for marketing approval.

For further information visit our website at www.pacifictherapeutics.com or email us at doug.unwin@pacifictherapeutics.com

Douglas H. Unwin, CEO & President

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FORWARD LOOKING STATEMENTS

Certain statements included in this press release constitute forward-looking information or statements (collectively, “forward-looking statements”), including those identified by the expressions “anticipate”, “believe”, “plan”, “estimate”, “expect”, “intend”, “may”, “should” and similar expressions to the extent they relate to the Company or its management. The forward-looking statements are not historical facts but reflect current expectations regarding future results or events. This press release contains forward looking statements. These forward-looking statements are based on current expectations and various estimates, factors and assumptions and involve known and unknown risks, uncertainties and other factors.

Readers should not place undue reliance on the Company’s forward-looking statements, as the Company’s actual results, performance or achievements may differ materially from any future results, performance or achievements expressed or implied by such forward-looking statements if known or unknown risks, uncertainties or other factors affect the Company’s business, or if the Company’s estimates or assumptions prove inaccurate. Therefore, the Company cannot provide any assurance that such forward-looking statements will materialize. The Company does not undertake to update any forward-looking information, except as, and to the extent required by, applicable securities laws.