CERTIFICATION INTERNATIONAL

Certificate of Registration

This is to certify that

Philippine Drug Enforcement Agency

PDEA Building, NIA Northside Road, National Government Center, Barangay Pinyahan, Quezon City, Philippines

operates a quality management system which has been assessed as conforming to

ISO 9001:2008

for the scope of activities

Provision of compliance and laboratory services.

Certificate No: **CIP/5065/15/02/913** Issue Date: **18 March 2015**
Valid until **17 March 2018** subject to adherence to the agreed ongoing audit programme, successful endorsement of certification following each audit and compliance with CI Regulations

Signed for and on behalf of

Managing Director

CERTIFICATION

CIP/5065

18 March 2015

UNDERSECRETARY ARTURO G. CACDAC, JR.
Director General
Philippine Drug Enforcement Agency
PDEA Building, NIA Northside Road,
National Government Center,
Barangay Pinyahan, Quezon City

Through: Engr. Salome DR. Jose
Quality Management Representative

Dear Undersecretary Cacdac,

Certification to ISO 9001:2008

After your successful completion of the requirements for certification, Certification International Philippines, Inc. hereby awards the *Philippine Drug Enforcement Agency* a Certificate of Registration to ISO 9001:2008. The Certificate is valid from 18 March 2015, with the following scope:

"Provision of compliance and laboratory services."

Your ISO 9001:2008 Certificate is valid for three years until 17 March 2018, subject to satisfactory results of semi-annual surveillance audits. The extension of your Certificate after this three-year period will be based on your effective conformity to the requirements of ISO 9001:2008, as determined by a re-assessment of your Quality Management System in its entirety. Please see attached surveillance audit schedule.

As a certified organization, you are required to maintain your Quality Management System to ISO 9001:2008 and to comply with CI's Conditions of Certification, including the use of CI's Certificate of Registration and Quality Mark in accordance with CI Regulations Part 4 and Part 5 (copy enclosed).

The validation of the consistent implementation and effectiveness of your improvement actions on the non-conformities cited during the Stage 2 audit will be carried out on the first surveillance audit.

Congratulations on your accomplishments with the use of ISO 9001:2008 as an instrument for enhancing the competitiveness of your organization and for meeting your customers' needs and expectations. We look forward to a strong partnership that will maximize your organization's benefits from its certified Quality Management System.

Best wishes for your continued success.

Very truly yours,
CERTIFICATION INTERNATIONAL

RENATO V. NAVARRETE
Managing Director

Encl: CI Regulation Parts 4 and 5