

Département de psychiatrie Centre de neurosciences psychiatriques Site de Cery CH-1008 Prilly - Lausanne

# Centre de Neurosciences Psychiatriques CNP SEMINAR

## ANNOUNCEMENT

## Friday, September 16, 2016, 11 a.m.

## "Past, Present and Future of Pharmacogenetic Testing in Psychiatry"

## Prof Jose de Leon

University of Kentucky Mental Health Research Center at Eastern State Hospital,

Lexington, Kentucky, USA

Invited by Chin Eap (Chin.Eap@chuv.ch)

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United States (US) agencies, particularly the Food and Drug Administration (FDA), have been at the forefront in attempting to make pharmacogenetic testing a reality. Therefore, understanding US regulations and practices is relevant in other countries.

Past and present pharmacogenetic testing in psychiatry in the US can be summarized in 3 phases: a) the fear phase (early 2000s), when pharmaceutical companies were afraid of pharmacogenetic testing; b) the failure phase (late 2000s), when the first commercialized pharmacogenetic tests failed and c) the hype phase (currently) when non-validated commercialized pharmacogenetic tests are aggressively marketed.

Regarding the present, the author's view is that pharmacogenetic testing can help in a few psychiatric cases: 1) before starting: a) carbamazepine in patients with Asian ancestry (HLA-B\*15:02 testing), or b) tricyclic antidepressants (CYP2D6 and CYP2C19 genotyping); and 2) occasionally when seeing lack of efficacy or adverse drug reactions with selective serotonin reuptake inhibitors, venlafaxine, pimozide, aripiprazole, brexpiprazole, iloperidone, risperidone, clozapine, or atomoxetine (CYP2D6 and/or CYP2C19 genotyping). Commercial tests should not be ordered for: 1) CYP1A2, CYP2B6, CYP3A4 or CYP3A5 genotyping; 2) brain neurotransmitter and/or transporter genotyping; or 3) diagnosing schizophrenia, depression or bipolar disorder.

Regarding the future, US marketing of pharmacogenetic tests requires: 1) understanding the pharmacological complexity of drug response, 2) modifying the oversight of non-FDA regulatory agencies, 3) clarifying the FDA's role, and 4) promoting innovative marketing. The incorporation of pharmacogenetic tests into long-term practice requires: 1) not jeopardizing pharmacogenetic testing by short-sighted marketing of non-validated tests, 2) educating prescribers about benefits, 3) educating patients about limitations, and 4) considering the differences between isolated testing and generalized testing incorporating big data.

### Selected Publications:

- 1) de Leon J. Pharmacogenetic tests in psychiatry: From fear to failure to hype (editorial). <u>Journal of Clinical</u> <u>Psychopharmacology</u> 36(4):299-304, 2016.
- 2) Spina E, de Leon J. Clinical applications of CYP genotyping in psychiatry. <u>Journal of Neural Transmission</u>122(1):5-28, 2015.



) de Leon J, Spina E. What is needed to incorporate clinical pharmacogenetic tests into the practice of psychopharmacotherapy? <u>Expert Review in Clinical Pharmacology</u> 9(3):351-4, 2016.