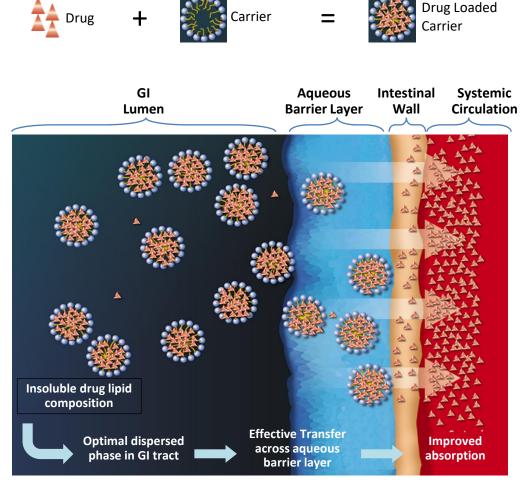
LPCN 1111

Once Daily Oral Testosterone



Drug Delivery Technology Leader with Specialty Pharma Strategy Initially Focused on Neuroendocrine and Metabolic Disorders



Giliyar et al. Drug Delivery Technology, Jan 2006, Vol 6 No.1



Lip'ral Drug Delivery Technology Platform

- 1) Invented by Renowned Formulation Experts
- 2) Validated Through a Recent FDA Drug Approval



3) Subject to Multiple Prior Partnerships











4) Protected by Robust WW Patent Portfolio >30 Issued & Pending Patents

Lipocine Pipeline





LPCN 1111: First Oral Once-A-Day "Phase 3 Ready" Candidate

Unique Prodrug of Testosterone



Convenient option for large and growing market



Patients and physicians preferred once a day oral testosterone



Positive Phase 2b study results



Strong IP Coverage



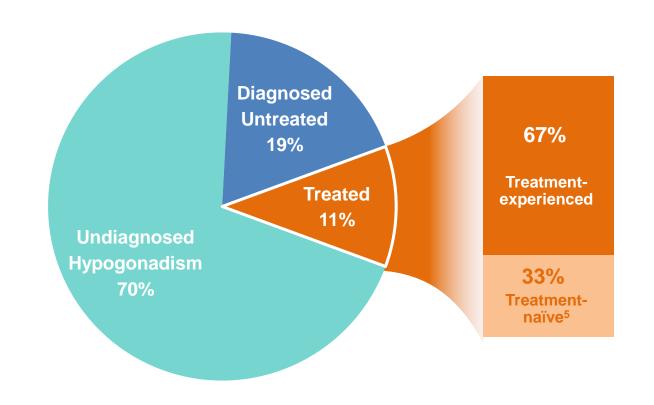
Hypogonadism Affects Up to 20M Men^{1, 2} in the United States

TLANDO Franchise has the Potential to Drive Market Expansion^{1,2}

Hypogonadism Under Treatment in US

Close to **6M** men with diagnosed hypogonadism³

2M men being treated⁴





Issues with Current TRT Options

Potential Barrier To Newly Diagnosed and Existing Patients











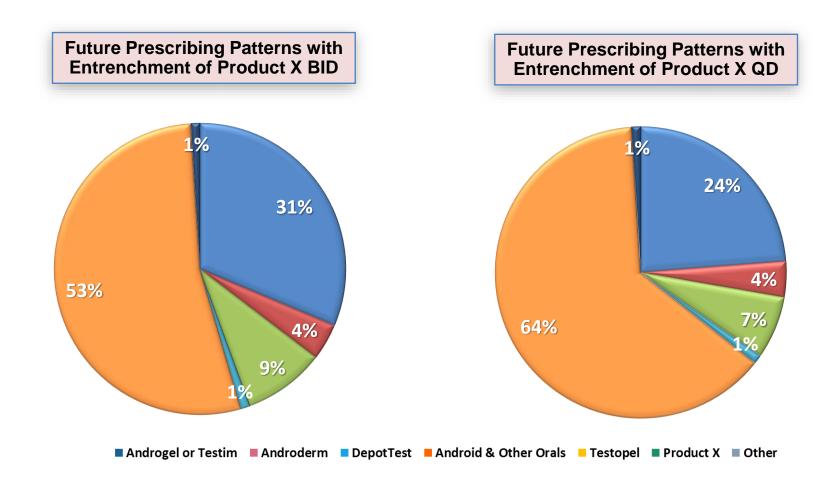


- Black box warning
 - Secondary exposure to testosterone
 - Pulmonary oil micro embolism (POME) and anaphylaxis shock
- Inconvenient application or painful injection
- High incidence of erythrocytosis was reported in patients on non-oral
 - 67% on injectable T, 35% on pellets*
- Most require dose titrations Majority of patients not on efficacious dose at start of therapy and require up titration
 - Avg titration time to reach efficacious dose takes 3-6 months
 - Finding appropriate dose through titration is burdensome for physicians and patients
 - Requires additional clinic/pharmacy visits and invasive samplings
- Poor persistence with products requiring titration
 - 50-80% drop off in 3-6 months
 - Insufficient T levels at the beginning of therapy is one of the top reasons patients stop using TRT



TLANDO XR: Market Research

Physician Intent-to-Prescribe Statistically Higher¹ with a QD





Phase 2b Study: Result Summary

Met Primary and Secondary Endpoints

- Once daily dose for 14 days in an open label, multiple dose PK study in hypogonadal men (n=36 subjects)
 - ✓ Phase 3 dose identified
 - ✓ No Drug related SAEs
 - Drug Related AEs are Mild to Moderate

Next step: Finalizing pivotal study protocol with FDA

