Forward Looking Statements

The statements in this presentation that relate to future plans, events or performance are forward-looking statements, which are made pursuant to the safe harbor provisions of Section 27A of the Securities Act of 1933, as amended. Such forward-looking statements involve significant risks and uncertainties, and actual results, events and performance may differ materially from those expressed or implied in this presentation. We have based these forward-looking statements largely on our current expectations and projections about future events and financial trends that we believe may affect our financial condition, results of operations, business strategy and financial needs. These forward-looking statements include, but are not limited to, statements concerning the following: (i) estimates of our expenses, future revenue and capital requirements; (ii) our ability to obtain additional funding on terms acceptable to us, or at all; (iii) our ability to advance product candidates into, and successfully complete, clinical trials; (iv) the initiation, timing, progress and results of future preclinical studies and clinical trials, and our research and development programs; (v) the timing or likelihood of regulatory filings and approvals; (vi) our expectations on sales and market acceptance for bremelanotide (Vyleesi®) for hypoactive sexual desire disorder (HSDD), a type of female sexual dysfunction (FSD), including our licensees outside North America jurisdictions; (vii) our expectation regarding timelines for development of our other product candidates; (viii) the potential for commercialization of our other product candidates, if approved for commercial use; (ix) our ability and the ability of our licensees to compete with other products and technologies similar to our product candidates; (x) the ability of third party collaborators to timely carry out their duties under their agreements with us and our licensees; (xi) the ability of contract manufacturers to perform their manufacturing activities in compliance with applicable regulations; (xii) our ability to recognize the potential value of our licensing arrangements with third parties; (xiii) the potential to achieve revenues from the sale of our product candidates; (xiv) our ability to maintain product liability insurance at a reasonable cost or in sufficient amounts, if at all; (xv) the retention of key management, employees and third-party contractors; (xvi) the scope of protection we are able to establish and maintain for intellectual property rights covering our product candidates and technology; (xvii) our compliance with federal and state laws and regulations; (xviii) the timing and costs associated with obtaining regulatory approval for our product candidates; (xix) the impact of legislative or regulatory healthcare reforms in the United States; and (xx) other risks disclosed in our SEC filings. The forward-looking statements in this presentation do not constitute guarantees of future performance. We undertake no obligation to publicly update these forward-looking statements to reflect events or circumstances that occur after the date of this presentation.
Pioneering a novel mechanism and approach to treating inflammatory & autoimmune diseases with a focus on ocular indications

- Demonstrated expertise moving programs from discovery to FDA approval
- Expertise in the biology and chemistry of the melanocortin system
- First company to procure FDA approval for a melanocortin agent (Vyleesi®)
- Strategy leverages our chemistry and biology across multiple therapeutic opportunities
- MOAs with the potential to modify underlying disease pathologies - not just treat symptoms
### Pipeline Development Programs

<table>
<thead>
<tr>
<th>Melanocortin Receptor Programs</th>
<th>Pre-clinical</th>
<th>Phase 1</th>
<th>Phase 2</th>
<th>Phase 3</th>
<th>NDA Submission</th>
<th>FDA Approval</th>
<th>Status/Next Steps</th>
</tr>
</thead>
<tbody>
<tr>
<td>PL9643 MCr Agonist Dry Eye Disease</td>
<td></td>
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<td></td>
<td></td>
<td>Phase 2 Dry Eye Trial Started 1Q2020 Positive Data 4Q2020 Phase 3 Trial to be Initiated 4Q2021 Phase 3 Data Expected 1H/2H2022</td>
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<td>PL9643 MCr Agonist Second Front of the Eye Indication</td>
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<td>Evaluating Several Indications Trial Initiation Targeted for 1H2022</td>
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<tr>
<td>MCr Agonist Diabetic Retinopathy Non-Infectious Uveitis</td>
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<td></td>
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<td></td>
<td>IVT Formulation Under Development</td>
</tr>
<tr>
<td>PL8177 MC1r Agonist (Oral) Inflammatory Bowel Disease</td>
<td></td>
<td></td>
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<td></td>
<td></td>
<td>Phase 2 Trial for Ulcerative Colitis to be Initiated 1H2022 with Data 2H2022</td>
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</table>

<table>
<thead>
<tr>
<th>Natriuretic Peptide Receptor Programs</th>
<th>Pre-clinical</th>
<th>Phase 1</th>
<th>Phase 2</th>
<th>Phase 3</th>
<th>NDA Submission</th>
<th>FDA Approval</th>
<th>Status/Next Steps</th>
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<tbody>
<tr>
<td>PL3994 NPR-A Cardiovascular Disease</td>
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<td></td>
<td></td>
<td></td>
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<td>Entered Phase 2a Clinical Trial Supported by Grant from the American Heart Association in 2H2020</td>
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<tr>
<td>PL5028 NPR-A/C Agonist Cardiovascular and Fibrotic Diseases</td>
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<td>Completed Preclinical Work Evaluating Options</td>
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</tbody>
</table>

### Commercial Product

| Vyleesi® (bremelanotide) Hypoactive Sexual Desire Disorder | FDA Approval 2Q2019 | Seeking U.S. and ROW Licenses |

Source: Company Filings
## Milestones

<table>
<thead>
<tr>
<th>Vyleesi (bremelanotide) for Hypoactive Sexual Desire Disorder</th>
<th>Status/Completion</th>
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</thead>
<tbody>
<tr>
<td>North American Rights Regained</td>
<td>3Q2020</td>
</tr>
<tr>
<td>China Licensee PK Study / S. Korea Licensee PK Study</td>
<td>2H2021</td>
</tr>
<tr>
<td>Seeking U.S. and ROW Partnerships</td>
<td>2H2021-2022</td>
</tr>
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</table>

<table>
<thead>
<tr>
<th>Melanocortin System Inflammatory &amp; Autoimmune Disease Programs</th>
<th>Initiated</th>
<th>Data</th>
</tr>
</thead>
<tbody>
<tr>
<td>PL9643 – Dry Eye</td>
<td></td>
<td></td>
</tr>
<tr>
<td>IND</td>
<td>4Q2019</td>
<td></td>
</tr>
<tr>
<td>Phase 2</td>
<td>1Q2020</td>
<td>4Q2020</td>
</tr>
<tr>
<td>Phase 3 (Melody 1)</td>
<td>4Q2021</td>
<td>1H/2H2022</td>
</tr>
<tr>
<td>Phase 3 (Melody 2 &amp; 3)</td>
<td>1H/2H2022</td>
<td>2H2023</td>
</tr>
<tr>
<td>PL8177 – Ulcerative Colitis</td>
<td>1H2022</td>
<td>2H2022</td>
</tr>
<tr>
<td>Phase 2 Ulcerative Colitis Proof-of-Concept</td>
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</tr>
<tr>
<td>PL9643-2nd Front of Eye Indication</td>
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<tr>
<td>Phase 2 Proof-of-Concept</td>
<td>1H2022</td>
<td>2H2022</td>
</tr>
<tr>
<td>MCR Agonist-Retinal indication</td>
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<tr>
<td>Proof-of-Mechanism DR/DME</td>
<td>2022</td>
<td>2023</td>
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<table>
<thead>
<tr>
<th>Natriuretic Peptide System Cardiovascular &amp; Fibrosis Programs</th>
<th>Initiated</th>
<th>Data</th>
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</thead>
<tbody>
<tr>
<td>PL3994 – Heart Failure</td>
<td>2H2020</td>
<td>2H2022</td>
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<tr>
<td>Open Label Phase 2a HF-pEF Patients</td>
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</table>
Melanocortin Inflammatory & Autoimmune Disease Programs
Pioneering a new way to patients

PALATIN
Immunological Effects of Melanocortin System

• Melanocortin system is up‐regulated by and integral to the resolution of autoimmune pathologies

• Modulates the activity of cells of the immune system

• Activated during disease state

• Activates resolution of proinflammatory processes
  ◦ Reduces NF-κB and other pro-inflammatory cytokines (IL-1, IL-2, IL-4, IL-6, IL-13, TNF-α, IFN-γ)
  ◦ Increased production of IL-10, an anti-inflammatory cytokine
  ◦ Mediates antigen specific T-cell and macrophage responses from pro-inflammatory to regulatory

• MC1R specific peptides and small molecules have demonstrated in vivo activity in numerous disease models of inflammation
Mechanism of Action of Melanocortins

Mechanism of Melanocortin Signaling, Leading to Inhibition of Inflammation

- Melanocortin receptor activates adenyl cyclase which generates cAMP activating protein kinase A (PKA)
- PKA activation leads to an influx of extravascular calcium and activation of IP3
- IP3 activates MAPK and JAK/STAT pathways, which inhibit degradation of cAMP and activate CREB. CREB prevents NFκB from activating transcription of inflammatory cytokines
- CREB is involved in downstream regulation of inflammation through production of immune signaling cytokines such as IL-10, which regulates the stimulation of immune cells

Immune cell

Gene transcription

Transcription factor

Calcium influx

Calcium

Calmodulin

CaMK-kinase

MAPK pathway

IP3

MEK2

ERK

PKA

cAMP

NFκB

B-RAF

MCC

PLC

Adenyl cyclase

Immune signaling cytokines

Nucleus
Melanocortin Anti-inflammatory Program

Rational design and synthesis of selective MC1R & MC1/5R agonists

PL8177: cyclic peptide selective MC1R agonist
PL9643: cyclic peptide MCR agonist

Reversal of pathology in multiple inflammatory and autoimmune disease models

Including inflammatory bowel disease, dry eye, uveitis, diabetic retinopathy & nephritis and pulmonary fibrosis

PL8177 Phase 2 clinical development candidate for indications requiring local or systemic administration

Oral formulation: Phase 2 for ulcerative colitis FPI 2H2021
Preclinical data in bleomycin pulmonary fibrosis model

Multiple opportunities for ocular indications

PL9643 topical DED
- Positive Phase 2 data
- Progressing to registrational studies
PL9643 topical for 2nd front of eye indication
PL8177 SC non-infectious uveitis Orphan designation
Candidate in IND-enabling activates for retinal indications
Ophthalmic Diseases: Front to Back

Conjunctiva
- Seasonal allergic
- Vernal (Orphan)
- Atopic (Orphan)

Conjunctiva/Cornea/Ocular surface
- Dry eye

Corneal Epithelium
- Toxicity from chemotherapy (Orphan)

Cornea endothelium
- Protect donor corneas for transplantation
- Improve corneal transplant survival
- Protection of cornea with cataract surgery

Ciliary Body
- Glaucoma therapy

Iris/Ciliary Body/Choroid
- Non-infectious uveitis

Retina
- Diabetic retinopathy
- Age-related macular degeneration

Optic nerve
- Neuro-protection in glaucoma
<table>
<thead>
<tr>
<th>REGION OF THE EYE</th>
<th>INDICATION</th>
<th>UNMET NEED</th>
</tr>
</thead>
<tbody>
<tr>
<td>Conjunctiva</td>
<td>Seasonal Allergic</td>
<td>Minimal unmet need</td>
</tr>
<tr>
<td></td>
<td>Vernal (orphan)</td>
<td>One product approved</td>
</tr>
<tr>
<td></td>
<td>Atopic (orphan)</td>
<td>Need for steroid-sparing agent</td>
</tr>
<tr>
<td>Conjunctiva/Cornea/Ocular surface</td>
<td>Dry eye</td>
<td>Huge market, tolerability is lacking</td>
</tr>
<tr>
<td>Corneal Epithelium</td>
<td>Toxicity from chemotherapy (Orphan)</td>
<td>Life-saving</td>
</tr>
<tr>
<td>Cornea endothelium</td>
<td>Protect donor corneas for transplantation</td>
<td>Unique indication</td>
</tr>
<tr>
<td></td>
<td>Improve corneal transplant survival</td>
<td>Unique indication</td>
</tr>
<tr>
<td></td>
<td>Protection of cornea with cataract surgery</td>
<td>Huge market, no therapies exist</td>
</tr>
<tr>
<td>Ciliary Body</td>
<td>Glaucoma therapy</td>
<td>Huge market, demand for new class of drugs, safety profile</td>
</tr>
</tbody>
</table>
## Ophthalmic Diseases: Unmet Medical Needs

<table>
<thead>
<tr>
<th>REGION OF THE EYE</th>
<th>INDICATION</th>
<th>UNMET NEED</th>
</tr>
</thead>
<tbody>
<tr>
<td>Iris/Ciliary Body/Choroid</td>
<td>Non-Infectious Uveitis</td>
<td>Need steroid-sparing therapy</td>
</tr>
<tr>
<td>Retina</td>
<td>Diabetic retinopathy</td>
<td>Largest market</td>
</tr>
<tr>
<td></td>
<td>Age-related macular degeneration</td>
<td>Largest market</td>
</tr>
<tr>
<td>Optic nerve</td>
<td>Neuro-protection in glaucoma</td>
<td>Unique</td>
</tr>
</tbody>
</table>
Dry Eye Overview

- Dry eye disease (DED) or keratoconjunctivitis is a multifactorial disorder of the tears and ocular surface
- Symptoms include dryness, irritation, redness, discharge and blurred vision
- Inflammation plays a prominent role in the development and amplification of the signs and symptoms of DED
- Current Treatments >$2 billion in revenue
  - Restasis®-topical cyclosporin
  - Xiidra®-topical integrin inhibitor
  - Topical steroids
  - Artificial tears
- Current treatments have efficacy and tolerability issues and there remains a high medical need for new innovative treatments that affect underlying disease processes
DED - One of the Most Common Eye Diseases in the United States\textsuperscript{1}

AMERICAN ADULTS suffer from symptoms of DED\textsuperscript{2,3}

29 million

= 1 out of every 7 American Adults\textsuperscript{2,3}

PL9643 Dry Eye Program

• PL9643 represents a novel approach to treating Dry Eye Disease (DED) by targeting the ability of the MS to resolve pathological inflammation
  ◦ PL9643 treats inflammation underlying the development and maintenance of DED, addressing both signs and symptoms of DED
• PL9643 is an agonist at the melanocortin 1 receptor (MC1r) and melanocortin 5 receptor (MC5r)
• PL9643 base patent runs to 2041
• Preclinical, DED studies PL9643 significantly reduced corneal epithelial damage with effects similar to Restasis®, a comparator reference agent
• Phase 2 study completed 2020
  ◦ Positive study
  ◦ 1st evaluation of MS agonist in ocular inflammatory indication
PL9643 Dry Eye Phase 2 Strategy

- Multi-center phase 2 RCT comparing PL9643 to placebo in DED patients with a 12-week treatment period
- Co-primary end points, a sign and symptom of DED
  - Sign: improvement in inferior fluorescein staining at week 12
  - Symptom: ocular discomfort measured at week 12
- Phase 2 study was exploratory with evaluations of multiple sign and symptom end points, patient segments, and time points
- Multiple outcomes can support a successful phase 2 DED study
  - Statistical significance for both co-primary end points
  - Statistical significance inferior fluorescein staining and a secondary symptom end point
  - Statistical significance ocular discomfort and a secondary sign end point
- Phase 3 registrational studies will need to achieve statistical significance on co-primary end points of a sign and symptom of DED
PL9643 Dry Eye Phase 2 Results

• Met primary objective of providing data required to advance into registration studies

• Statistical significance for the primary endpoints was not achieved in the ITT population that included mild, moderate, and severe patients

• In the sub-population of moderate to severe patients (N=61), PL9643 achieved statistical significance (P value <0.05 vs. vehicle) at week 2 and week 12 for multiple signs and symptoms

• PL9643 demonstrated excellent ocular safety and tolerability
  ◦ No drug related serious adverse or adverse events
  ◦ No drug related discontinuations
  ◦ High ocular comfort

• Favorable emerging product profile
  ◦ Rapid onset, well tolerated and global efficacy
Difference between PL9643 & Placebo for Corneal Fluorescein Staining Patients with Moderate or Severe Disease

In patients with moderate or severe disease significant improvement was observed for PL9643 compared with placebo for the primary endpoint of inferior corneal fluorescein staining (P<0.05)
Differences between PL9643 & Placebo in Conjunctival Redness and Tear Film Break-up Time
Patients with Moderate or Severe Disease

Signs of conjunctival redness showed numeric improvements (as demonstrated by negative change from baseline) and tear film break-up time showed significant improvement at 12 weeks.
Differences between PL9643 & Placebo Ora Calibra® Ocular Discomfort & 4-Symptom Questionnaire† Score's patients with Moderate or Severe Disease

PL9643 demonstrated significant improvement in ocular discomfort over placebo at Week 2 (as shown by negative change from baseline)

†Measured on 0-5 continuous scale
PL9643 Dry Eye Commercial Opportunity

DED is estimated to affect over 20 million people in the United States
- Majority of people suffering from DED are in the moderate to severe category (>75%)
- Most patients have persistent disease (>5yrs)

Existing therapies for dry eye disease generally regarded as inadequate by many physicians and patients
- Limited clinical trial evidence for both signs and symptoms
- Require weeks or months to demonstrate activity
- High discontinuation due to slow onset, lack of efficacy and high rates of side effects

PL9643 has a favorable commercial product profile / Differentiating factors to current approved therapies
- Quick onset of efficacy
- Excellent tolerability profile
PL9643 Dry Eye Disease Development – Status/Next Steps

- **Submitted Phase 2 Study to FDA**
  - 1H2021

- **EOP2 FDA Meeting Held**
  - 1H2021

- **Completed Non-clinical Activities**
  - 1H2021

- **Initiate Phase 3 Study**
  - 2H2021
  - Data 1H/2H2022

- **Initiate 2nd Phase 3 Study**
  - 1H/2H2022

- **NDA to FDA Submission**
  - 2H2023
Diabetic Retinopathy & Macular Edema

- By 2050, the number of Americans with diabetic retinopathy is expected to nearly double, from 7.7 million to 14.6 million
- DME affects ~10% of people with diabetic retinopathy
  - ~750,000 in the USA & 2.2 million people in the EU
- IVT VEGF antagonists and steroids are the main treatments for DME
  - Annual global sales for DR/DME estimated at $1.85b
- There is a high need for additional treatments
  - To delay progression, maintain and improve visual acuity
  - Replacement for steroids without glaucoma or cataract side effects
- Our melanocortin agonist have been evaluated in multiple animal models of retinal disease
• PL8331 is a melanocortin receptor agonist
• VEGF and TNF-α levels are similar to healthy mice even though the diabetic mice remain hyperglycemic throughout the study
• IL-10 is a marker of inflammation resolution
Melanocortin Agonists Preserve Contrast Vision and Acuity

Rat diabetic retinopathy model
PL8177 Non-Infectious Uveitis

- Non-Infectious Uveitis (NIU) is a potentially blinding intraocular inflammatory disease that arises without a known infectious trigger and is often associated with immunological responses to unique retinal proteins

- Prevalence of NIU in N. America
  - Adults: ~72,000
  - Pediatric: ~21,000

- NIU causes bilateral legal blindness in 6% of patients and unilateral blindness in 18% of patients

- Only 2 FDA approved treatment options
  - Ozurdex® (dexamethasone intravitreal implant)
  - Humira® (adalimumab)
  - Significant off-label treatments – steroids, infliximab, methotrexate, azathioprine, etc.

- There remains a high need for new safer treatments
  - Use of steroids leads to glaucoma and cataracts and has systemic toxicities
  - Humira® increases serious infection risk and has substantial contraindications

- Orphan drug designation for PL8177 for NIU
PL8177 Experimental Autoimmune Uveitis

MC1R agonism has significant effects in reversing uveitis

Conducted in collaboration with Dr. A. Taylor at Boston University School of Medicine
Ocular Development Programs

- Demonstrated efficacy, safety and tolerability in phase 2 DED study
  - 1st evaluation of melanocortin system therapeutic in ocular inflammatory indication
  - Establishes translation of preclinical data into humans
  - Moving forward into registrational studies

- PL9643 phase 2 exploratory study in 2nd front of the eye indication

- PL8177 SC non-infectious uveitis Orphan Disease Designation ready for phase 2 proof-of-concept study

- MCR agonist demonstrated efficacy in animal models of retinal disease
  - Suppresses VEGF production and reduces vascular leakage
  - Preserves retinal structure
  - Suppresses inflammation and promotes resolution of inflammatory activity
  - Maintains visual acuity
PL8177 for Ulcerative Colitis
Why a Melanocortin Peptide for Ulcerative Colitis?

- MC1r is found on epithelial cells of the colon and appear to be accessible from the lumen of the colon
  - Evidence from preclinical animal studies
- PL8177 is a highly potent peptide of 7 aa that is a selective agonist at MC1r
- PL8177 main metabolite is PL8435 which maintains MC1r selective and agonist activity
- Most treatments for UC are systemic and have tolerability and safety limitations
- PL8177 is not systemically absorbed
  - Potential for excellent efficacy without safety concerns
- Compatible with Eudragit polymers that allow for oral delivery to the GI tract for topical delivery
Opportunity for PL8177 in UC Treatment Landscape

Severity: Mild | Moderate | Severe
---|---|---
Patient number: Large | | Small

**Remission induction**
- 5-ASA (Lialda)
- PL8177
- 5-ASA failure
- Steroids enema/suppository
- Steroids oral
- Steroids infusion
- Biologics & Immuno-modulators (Adalimumab BS)
- Surgical operation

**Remission maintenance**
- 5-ASA (Lialda)
- PL8177
- Biologics & Immuno-modulators (Adalimumab BS)

Drug price:
- Low
- High
PL8177 (20ug, 50ug and 100ug) and PL9680 (50ug) are given as a polymer formulation placed into #9 capsules.

Placebo capsules and test article capsules are administered twice daily.

Mesalazine is administered orally once daily.
Histological Findings: Total Colitis Index

- The scoring was based on examining three sections from each colon per animal:
  - Sections were taken at the distance of 2.5cm, 5cm and 7.5cm from the anus
  - Total colitis index includes observations
    - abnormalities of mucosal architecture,
    - extent of inflammation
    - erosion or ulceration
    - epithelial regeneration
    - percentage involvement by the disease process
PL8177 Oral Formulation Clinical Study 102 Summary

Confirmation of local colonic delivery based on presence of PL8177 and metabolite

- This was a phase 0, open-label clinical study conducted at a single center in The Netherlands designed to examine an oral formulation of PL8177
  - A microdose study was chosen in order to assess whether the oral formulation was delivered to the appropriate part of the gastrointestinal tract using a subclinical dose and very small amount of radioactivity

- The primary objectives were
  - To demonstrate release of [\(^{14}\text{C}\)]-PL8177 from the polymer-bound form of [\(^{14}\text{C}\)]-PL8177 in the colon after oral administration through observation of the main metabolite
  - To confirm that the orally administered, polymer-bound form of [\(^{14}\text{C}\)]-PL8177 did not result in systemic exposure to [\(^{14}\text{C}\)]-PL8177 and/or its [\(^{14}\text{C}\)]-PL8435 (main metabolite)
  - To establish the relationship between an oral dose of polymer-bound [\(^{14}\text{C}\)]-PL8177 and the amount of [\(^{14}\text{C}\)]-PL8177 and/or [\(^{14}\text{C}\)]-MAIN METABOLITE in the colon

- The secondary objective was to evaluate the safety and tolerability of the orally administered, polymer-bound form of [\(^{14}\text{C}\)]-PL8177 in healthy male subjects
PL8177 Oral Formulation – Human PK (microdose study)

Confirmation of GI tract release & no systemic exposure

Mean Percent Excretion vs Time in Feces of [14C]-PL8177 and [14C]-PL8435

Cohort 1 = [14C]-PL8177 + laxative 5 h postdose
Cohort 2 = [14C]-PL8177 + laxative 8 h postdose
Cohort 3 = [14C] PL8177 + laxative 11 h postdose
Cohort 4 = [14C]-PL8177 + laxative 14 h postdose
Cohort 5 = [14C]PL8177 + laxative 17 h postdose
Cohort 6 = [14C]-PL8177 without laxative

- The metabolite [14C]-PL8435 was quantifiable in fecal samples from 10 subjects, demonstrating that orally administered [14C]-PL8177 was released from its polymer-bound formulation and metabolized to [14C]-PL8435 in the GI tract in those subjects.

- Neither the parent drug [14C]-PL8177 nor the metabolite [14C]-PL8435 were quantifiable in any of the plasma or urine samples, suggesting that oral administration of the polymer-bound formulation of [14C]-PL8177 does not result in systemic exposure of PL8177 and PL8435.
PL8177 Oral Formulation for UC – Summary Status

• Multiple positive animal model proof of efficacy data in gold standard disease model

• Efficacy as good/better than 5-ASA and glucocorticoids in animal model data

• Phase 1 SAD/MAD study with the systemic formulation (SC) up to 3mg for 7 days, up to 5mg SC for a single dose

• Phase 1 radiolabeled micro-dose study with the oral formulation, confirmed colonic delivery of oral PL8177

• Orally dosed PL8177 remains in the colon – there is no systemic exposure

• Phase 2 proof-of-concept trial initiation 4Q2021 / Data 2H2022
Vyleesi Operations/ Performance

OBJECTIVE

Demonstrate the commercial value and upside of Vyleesi and re-license to a committed partner

• **Vyleesi is a valuable asset in the ‘right’ hands**
  o FDA approved product with limited competition

• **FSD market**
  o Significant awareness needed / greater HCP and patient engagement

• **For the quarter ended September 30, 2021**
  o Gross product sales increased 18%, net revenue increased 98%, net revenue per prescription dispensed increased 45%, despite a 13% decrease in total prescriptions dispensed, over the prior quarter ended June 30, 2021
  o Gross product sales amounted to $1.4 million, with net product revenue of $159,482, compared to gross product sales for the period July 25 (the date Palatin regained North American rights to Vyleesi) to September 30, 2020, of $809,100, with negative net product revenue of $(288,560)
  o Market access, reimbursement coverage, and refill rates increased over the prior quarter ended June 30, 2021 and the quarters ended March 31, 2021 and December 31, 2020

• **Learn more about HSDD and Vyleesi at** [www.vyleesi.com](http://www.vyleesi.com) and [www.vyleesipro.com](http://www.vyleesipro.com)
FDA Approved Vyleesi®

Helping Premenopausal Women with Hypoactive Sexual Desire Disorder (HSDD)

Hey, you. Meet Vyleesi.
...it’s Now Approved

Vyleesi® is the first and only as-needed® treatment for premenopausal women with acquired, generalized hypoactive sexual desire disorder (HSDD).

*Administer subcutaneously as needed at least 45 minutes before anticipated sexual activity. The duration of its effect after each dose is unknown. Do not administer more than one dose within 24 hours or more than 8 doses per month.
HSDD is a Significant Market Opportunity

Number of premenopausal women who have low desire with associated distress

1/10\(^{1,2}\)

Affects 5.8 million U.S. premenopausal women\(^3\)
(1 in 10 premenopausal women)\(^{1,2}\)

98% (5.7M) of affected premenopausal women not on therapy\(^3\)

Focused on relevant digital channels

Creating an online community for HSDD patients
- Provide accurate information
- Tools to support the HSDD patient - symptom check, speaking with your doctor and additional resources

Ensure HCP readiness, provide information and tools to diagnose and treat HSDD patients with Vyleesi

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\(^3\) Palatin supported research that was performed by Burke, Inc., an ISO 20252-certified company, in compliance with the established standard for market, opinion, and social research.
# Financial Snapshot

## Financial Highlights as of September 30, 2021

<table>
<thead>
<tr>
<th>Description</th>
<th>Amount</th>
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<tbody>
<tr>
<td>Cash and Cash Equivalents</td>
<td>$53.4 million</td>
</tr>
<tr>
<td>Accounts Receivable</td>
<td>$0.9 million</td>
</tr>
<tr>
<td>Inventory</td>
<td>$1.1 million</td>
</tr>
<tr>
<td>Inventory Purchase Commitments (over the next 5 years)</td>
<td>$9.8 million</td>
</tr>
</tbody>
</table>

## Summary Capitalization as of October 31, 2021

<table>
<thead>
<tr>
<th>Common Shares and Equivalent</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>231.3 million shares</td>
<td>Common Stock</td>
</tr>
<tr>
<td>0.1 million shares</td>
<td>Preferred</td>
</tr>
<tr>
<td>4.6 million shares</td>
<td>Warrants</td>
</tr>
<tr>
<td>21.9 million shares</td>
<td>Options</td>
</tr>
<tr>
<td>13.1 million shares</td>
<td>RSUs</td>
</tr>
<tr>
<td>271.0 million shares</td>
<td>Fully Diluted Shares</td>
</tr>
</tbody>
</table>
THANK YOU