



Innovators in Complex Carbohydrate Chemistry™

Corporate Presentation June 2013

OTCQB: BTHE
www.bostonti.com

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Company Description

A pharmaceutical company focused on the development of novel compounds based on complex carbohydrate chemistry to address unmet medical needs in diabetes and inflammatory disease



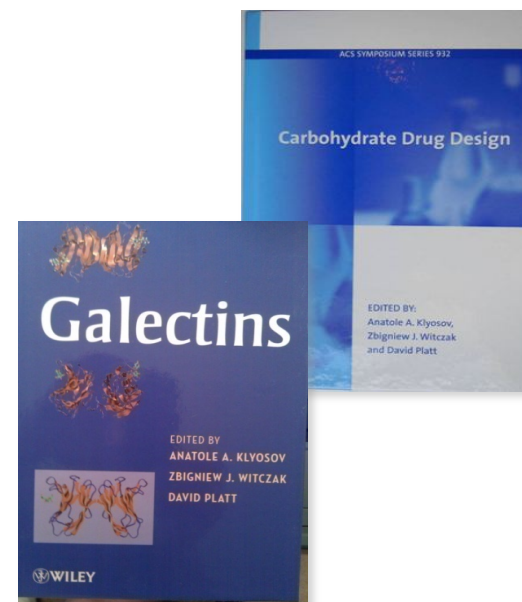
Investment Highlights

- Two products that address unmet needs in the large and growing diabetes drug market
- Multiple clinical and regulatory milestones over the next two years
- Lead drug provides a safe, non-systemic approach that works in combination with other drugs that meets a true market need
- Patented chemistry provides long-term high value assets
- Management leverages domain expertise in carbohydrate engineering to advance product development



Unique Chemistry Design

- Growing interest in carbohydrates as a therapeutic
 - Biological importance is now better understood
 - Plays fundamental role in normal cell functions
 - Participates in cell-cell interactions
 - Stimulates immune response
- David Platt, Ph.D. is an expert/pioneer in galectin research; inventor of many patents
- Founder & CEO of three publicly traded companies:
 - Pro-Pharmaceuticals now Galectin Therapeutics (Nasdaq: GALT) - Liver /cancer
 - SafeScience (now LaJolla Pharmaceuticals OTC: LJPC) - Kidney
 - Boston Therapeutics (OTCQB: BTHE) - Diabetes
- Co-editor of *Carbohydrate Drug Design and Galectins*
 - Influential volumes in the design of drugs using complex carbohydrates





Experienced Management Team

Team	Background
<p>David Platt ,Ph.D. Chief Executive Officer Chief Financial Officer Chairman</p>	<ul style="list-style-type: none">• 2001-2009: CEO/Chairman of Pro-Pharmaceuticals, Inc., now Galectin Therapeutics, Inc. (NASDAQ: GALT)• 1995-2000: CEO, Chairman and founder of SafeScience Inc., a Nasdaq-listed company• 1992-1995: CEO, Chairman and founder of International Gene Group, Inc., the predecessor company to SafeScience, Inc.• 1989-1991: Research fellow at the Michigan Foundation (now Barbara Ann Karmanos Institute)• Research fellow at the Weizmann Institute of Science, Rehovot, Israel• Ph.D. in Chemistry in 1988 from Hebrew University in Jerusalem• Published peer-reviewed articles and holds many patents, primarily in the field of carbohydrate chemistry
<p>Kenneth A. Tassej, Jr. President</p>	<ul style="list-style-type: none">• Co-founder and President of Boston Therapeutics, Inc. since November 2010• CEO & President of Boston Therapeutics from 2009 until merger with Avanyx Therapeutics• Former President of TKCI (consulting firm for commercial finance) from 2007 to 2009
<p>Jonathan B. Rome Chief Operating Officer</p>	<ul style="list-style-type: none">• More than 30 years executive experience within the pharmaceutical industry• Founder, President & CEO of ThePharmaNetwork, LLC from 2000 to 2012• Specialist in pharma portfolio development, licensing, sales, marketing and distribution of pharmaceuticals and active pharmaceutical ingredients
<p>Anthony Squeglia Director of Strategic Planning</p>	<ul style="list-style-type: none">• Former CFO for Galectin Therapeutics, Inc. (Nasdaq: GALT) and its predecessor company Pro-Pharmaceuticals, Inc. (OTC: PRWP) from 2007 to 2012• VP Investor Relations for Pro-Pharmaceuticals from 2003 to 2007• Senior management positions at Unisys, AT&T, Summa Four, Quentra Networks, Colonial Penn, ITT• BBA from Wharton, MBA from Pepperdine



Medical & Scientific Advisory Team

Team

Hana Chen-Walden, M.D.
Chief Medical Director

Dr. Peter Sheehan, M.D.
Advisor, Medical Director

Shih-Chun David Liu, Ph.D.
Scientific Advisor

Dr. Eliezer Zomer, Ph.D.
Scientific Advisor

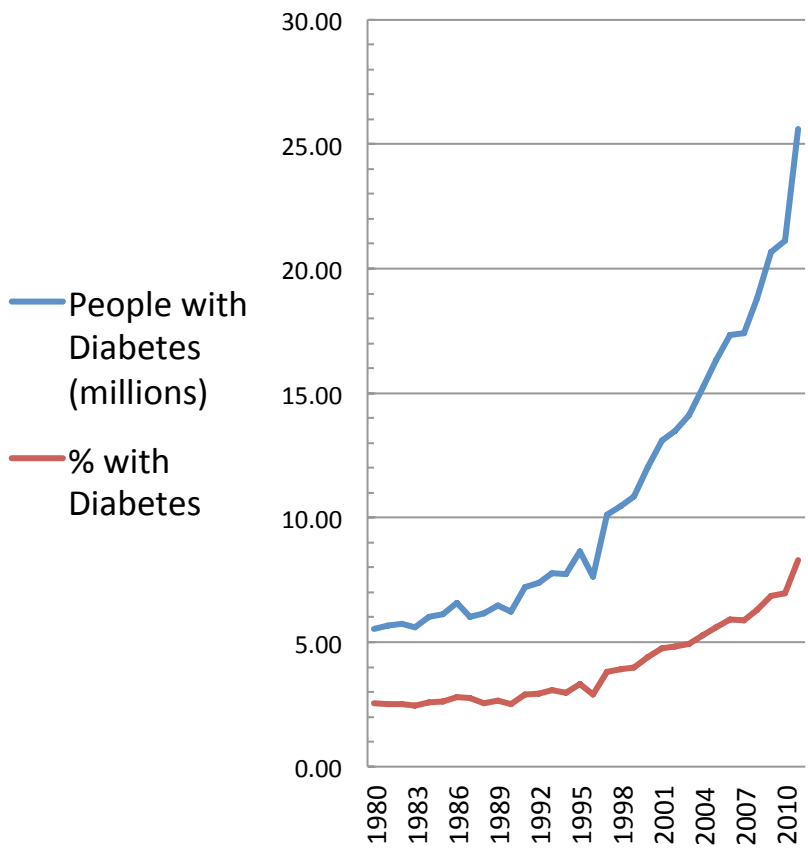
Background

- Specialist in regulatory affairs in the pharmaceutical industry in U.S. and Europe
 - 30 years of regulatory experience with the EMEA and in individual European countries
 - Consultant since 2004 for European Clinical and Regulatory Consultancy
 - M.D. from University of Tel Aviv, Israel
- American Diabetes Association:
 - Current President, NYC Leadership Council
 - Current Chairman of Cardiometabolic Risk Initiative
 - Former national Board Member
 - Internationally respected Endocrinologist and diabetes specialist
 - Clinical interest in peripheral artery disease, diabetic neuropathy, wound healing
- Research contributed to the fundamental understanding of the red blood cell membrane architecture and identified surface receptor for malaria invasion
 - Published over sixty research papers in prestigious journals
 - Former research scientist in the Biomedical Research Department at Tufts Medical School
 - Associate Professor of Medicine, Tufts University School of Medicine since 1992
 - Ph.D. in biochemistry from Carnegie-Mellon University
 - VP, founder at HDM Systems Corp., a green power electronic devices developer
- Executive VP of Manufacturing and Product Development of Galectin Therapeutics (GALT) (formerly Pro-Pharmaceuticals) since 2000
 - Founder of Alicon Biological Control
 - Former VP of product development at SafeScience, Inc. and former VP of R&D at Charm Sciences, Inc.
 - Ph.D. in biochemistry from the University of Massachusetts



Diabetes: A Growing Epidemic

US Population with Diagnosed Diabetes



Diabetes Facts (US)

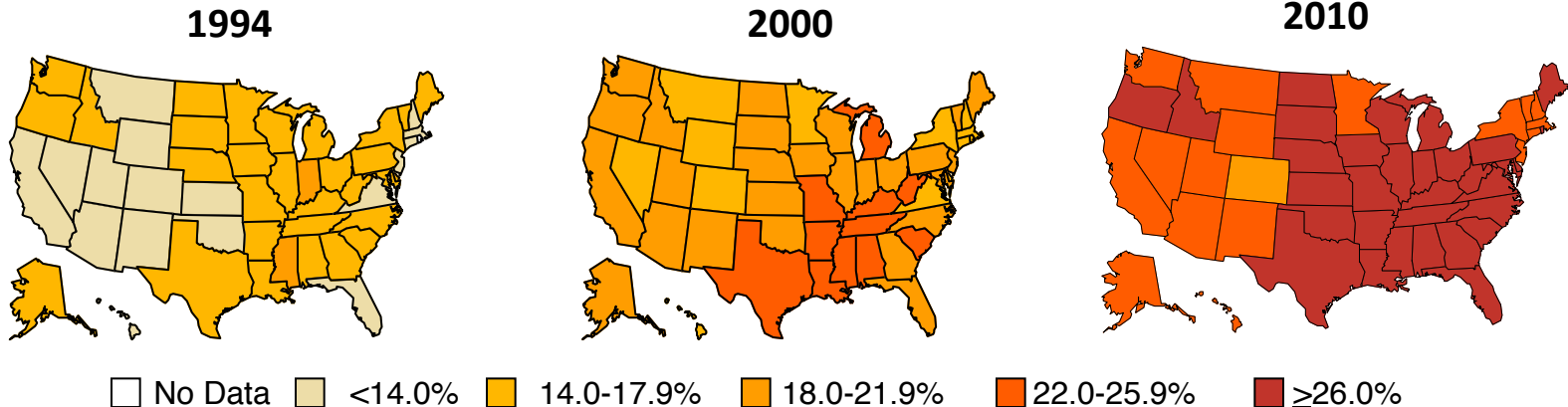
- 26 mm people with diabetes (8.3% of population)
 - 460% increase since 1980
- Additionally, 79 mm are pre-diabetic
- 1 of 3 US adults will have diabetes by 2050 if current trends continue
- Leading cause of:
 - kidney failure
 - non-traumatic lower-limb amputations
 - new cases of blindness
- Major cause of heart disease and stroke

Source: CDC Division of Diabetes Translation. National Diabetes Surveillance System and 2011 CDC Diabetes Fact Sheet

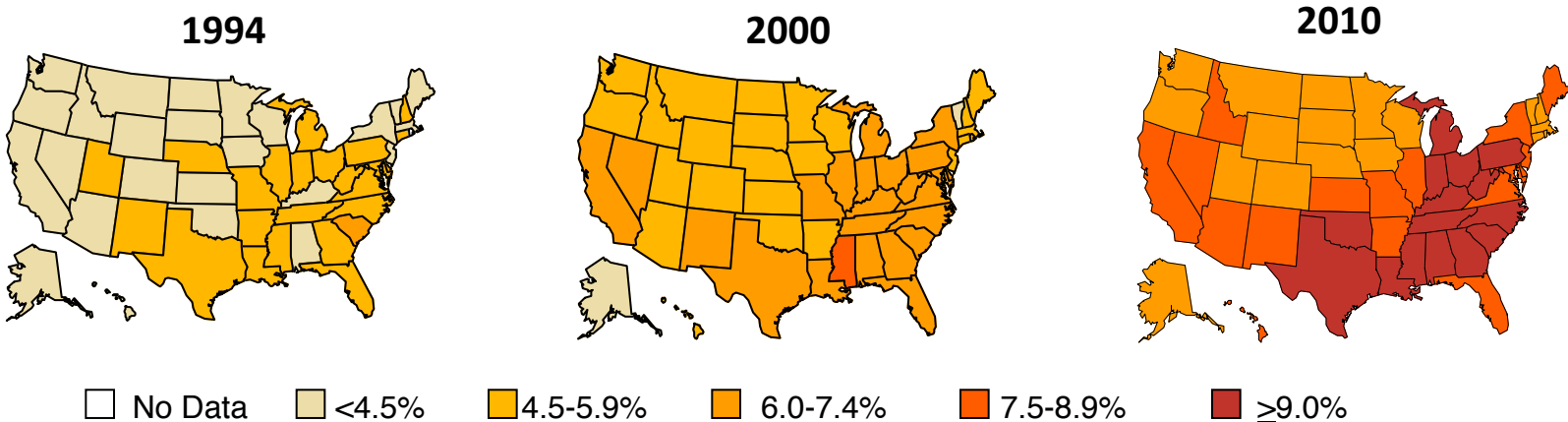


Diabetes: A Growing Epidemic

Obesity (BMI ≥30 kg/m²)



Diabetes



Source: CDC Division of Diabetes Translation. National Diabetes Surveillance

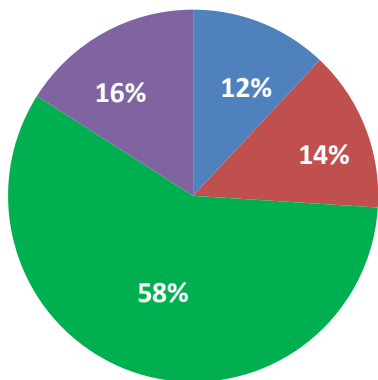


Economic Considerations in US

Diabetes effect on US Healthcare Dollars

- **\$245 billion:** Total estimated costs of diagnosed diabetes in 2012
- **↑ 10%** spent directly on diabetes and its complications
- **↑ 20%** spent on caring for people with diagnosed diabetes

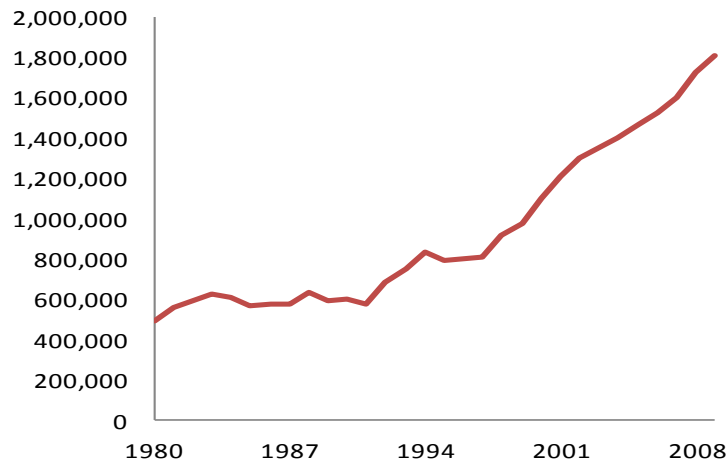
Treatment Type of Diagnosed Diabetes Patients in US Adults



Lead oral drug candidate shows safety when added to oral anti-diabetic medications and insulin in clinical study

- Insulin only
- Insulin and oral medication
- Oral medication only
- No medication

New Cases of Diagnosed Diabetes in US Adults Aged 18–79 Years

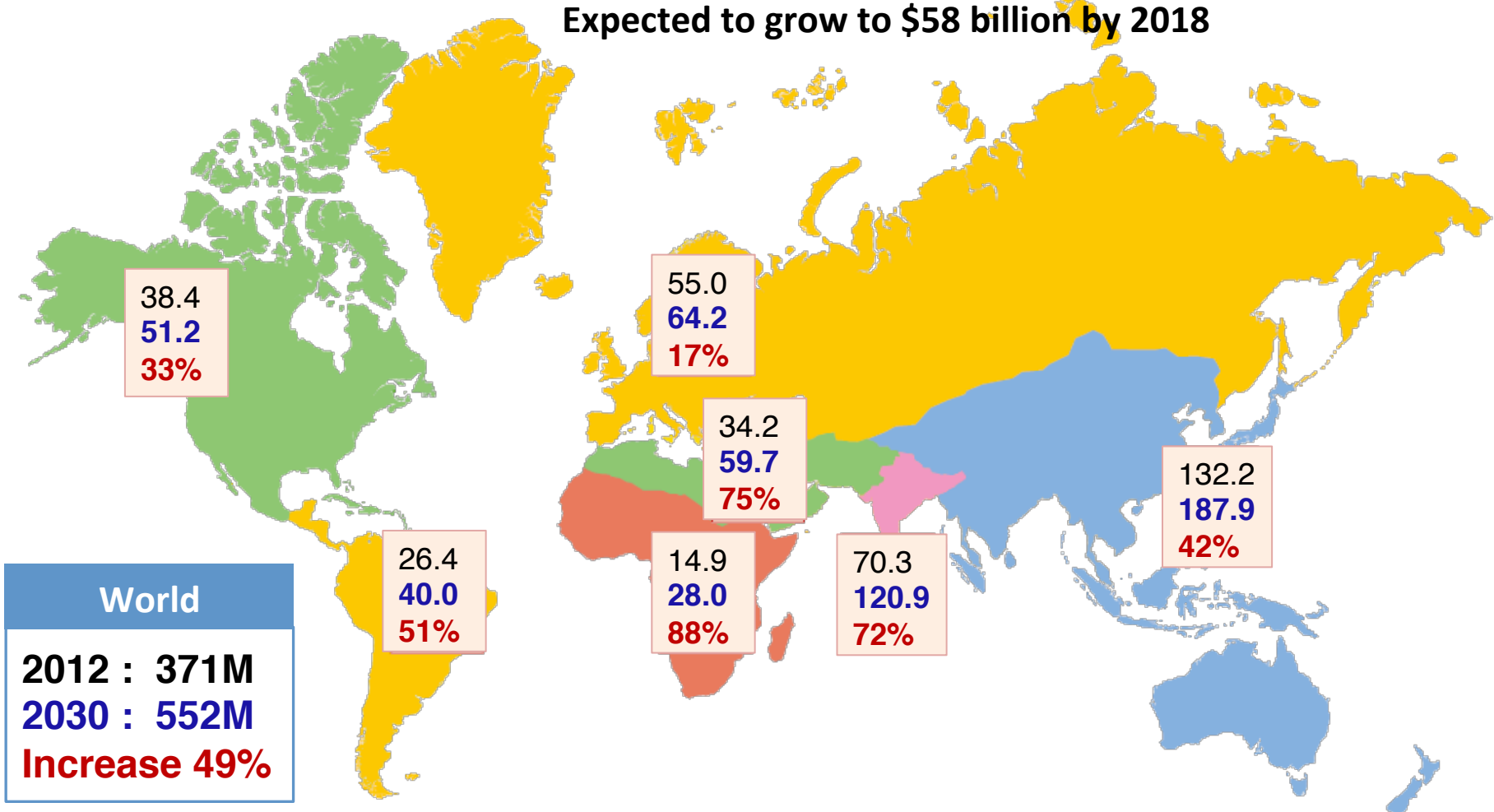


Sources: American Diabetes Association; Centers for Disease Control and Prevention, and Standard & Poor analyst report (Oct 04, 2012)



Diabetes Growth Projections 2012-2030

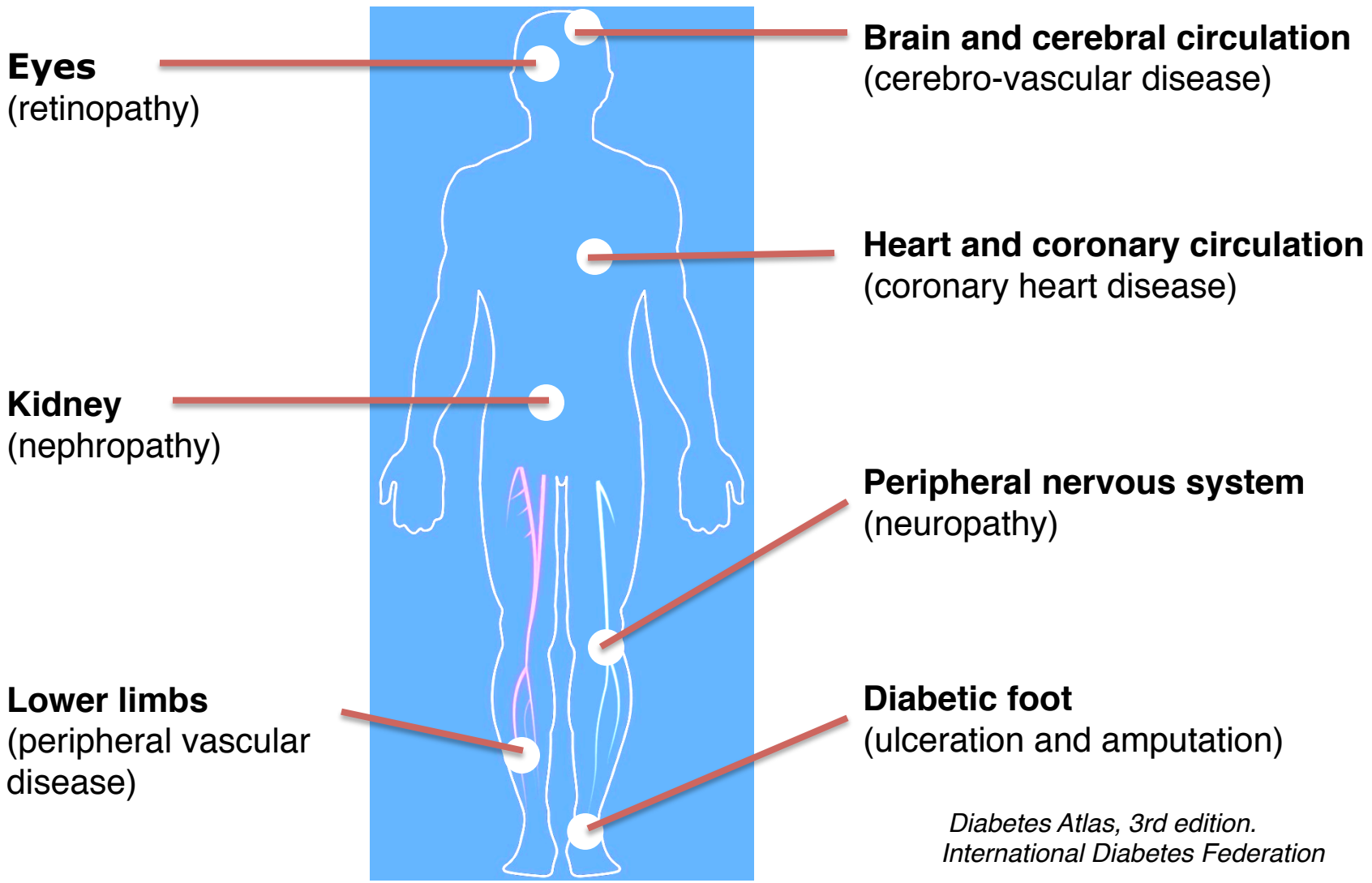
Diabetes worldwide drug market size \$35 billion
Expected to grow to \$58 billion by 2018



Source: International Diabetes Federation Diabetes Atlas 5th Edition: 2012 Update



Diabetes Complications



*Diabetes Atlas, 3rd edition.
International Diabetes Federation*



Product Pipeline

Product	Description	Indication	Preclinical	Phase I	Phase II	Phase III
PAZ320	Chewable tablet that manages blood sugar in combination with other treatments	Type 2 Diabetes				
IPOXYN	Injectable anti-necrosis drug	Lower-limb ischemia in diabetic patients				

- PAZ320 works in the gastrointestinal tract limiting side effects, unlike other diabetes treatments
- IPOXYN is an injectable anti-necrosis drug that treats hypoxia, which is the lack of oxygen in living cells



PAZ320

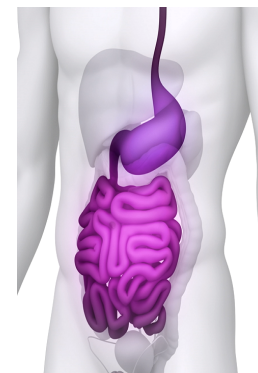
- Complex carbohydrate chemical structure
- Chewable drug taken before meals that prevents and treats Type 2 diabetes
- Novel, non-systemic approach to blood sugar management that works in combination with other drugs
- **Efficacy and safety in combination with metformin**
 - 50 million prescriptions in US per year
- Strong safety profile
 - No serious adverse events (SAE)
 - provides a competitive advantage compared to other anti-diabetic drugs

PAZ 320

Non-Systemic

PAZ 320 works locally in the gastrointestinal tract

Less risk for side effects



Other Diabetes Drugs

Systemic

Typical mechanisms involve interaction with liver, kidney, pancreas and cells



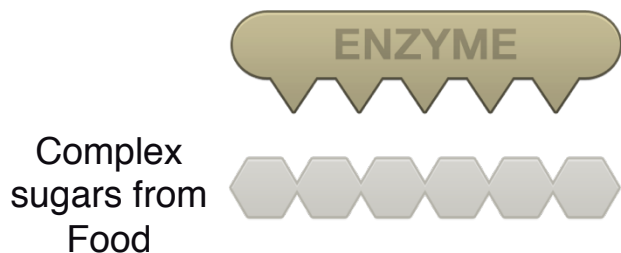


PAZ320: Mechanism of Action

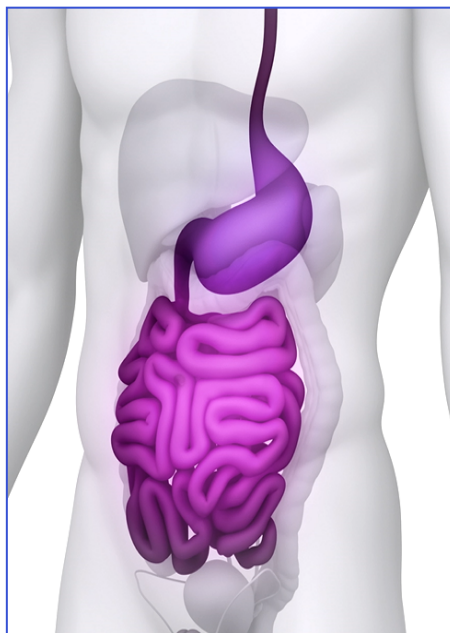
Carbohydrate-hydrolyzing enzyme inhibition
significant reduction in post-meal elevation of glucose

Without PAZ 320

Enzymes break down complex sugars into simple sugars

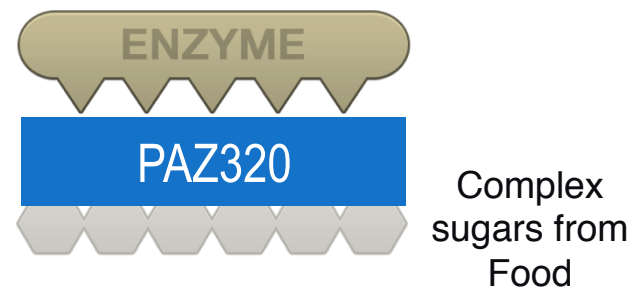


More Glucose Available for Absorption (BAD)



With PAZ 320

Inhibits enzymes that release glucose from complex carbohydrates



Less Glucose Available for Absorption (GOOD)



Diabetes Drugs: Side Effects

Drug Brand	Cardiac Events	Stroke	Cancer Risk	Eye Damage	Lactic Acidosis	Diarrhea	Nausea and Vomiting	Severe Stomach Pain	Risk of Bone Fracture	Pancreatitis	Hypoglycemia	Respir. Tract Infection	Risk of Bone Fracture	Urinary Tract Infection	Mild GI symptoms
PAZ320 (Prandiol)															✓
Precose (Acarbose)						✓	✓	✓							
ACTOS (Pioglitazone)	✓	▪	✓	✓	▪				✓	▪		▪	✓	▪	▪
Avandia (Rosiglitazone)	✓	✓	✓	✓	▪		▪	▪	✓	▪			✓	▪	▪
Bydureon (Exenatide)			✓			✓	✓	▪		✓	✓				
Byetta (Exenatide)	▪					✓	✓	▪		✓	✓				
Invokana (Canagliflozin)	✓	✓							▪			▪	▪	✓	▪
Januvia (Sitagliptin)	✓		✓						▪			✓	▪	▪	▪
Glucophage (Metformin)			▪		✓	✓	✓	▪		▪					
Onglyza (Saxagliptin)	✓		✓			✓			▪	✓	▪	✓	▪	▪	▪
Victoza (Liraglutide)			✓							▪	✓	✓			
Trajenta (Linagliptin)						✓				✓	▪				

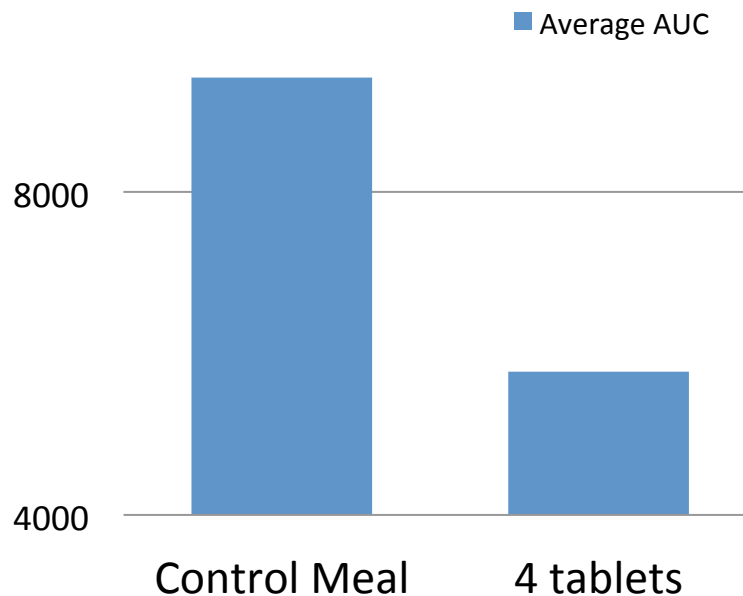
Sources: Medline Plus <http://www.nlm.nih.gov/medlineplus/>



PAZ320: Phase II Trial Results

- 45% of responders reported a 40% decrease in increased glucose post-meal
- No serious adverse events
- Sense of fullness was observed
- Works no matter how long the patient has had diabetes
- Works regardless of other medications

Responder Summary
Average Area Under the Curve
(p-value: ANOVA Friedman 0.012)



Trial conducted at Dartmouth Medical Center



PAZ320: Current and Planned Trials

Trial Type	Status	Patient Population	Goals
Phase II Study Dartmouth-Hitchcock Medical Center, US	Completed Data expect to be published in Endocrine Practice Q3 2013	<ul style="list-style-type: none">• 24 people with Type 2 diabetes• Currently using oral agents or insulin	<ul style="list-style-type: none">• Efficacy and safety• In combination with oral anti-diabetic medications and insulin
Phase II France	Initiating	<ul style="list-style-type: none">• 24 people with Type 2 diabetes currently using metformin	<ul style="list-style-type: none">• Efficacy and safety
Phase III US, Hong Kong, Korea and China	Planned Collaboration with major US diabetes clinic	<ul style="list-style-type: none">• 300 patients	<ul style="list-style-type: none">• Evaluation of the effects of PAZ320 with metformin on glucose AUC and HbA1c

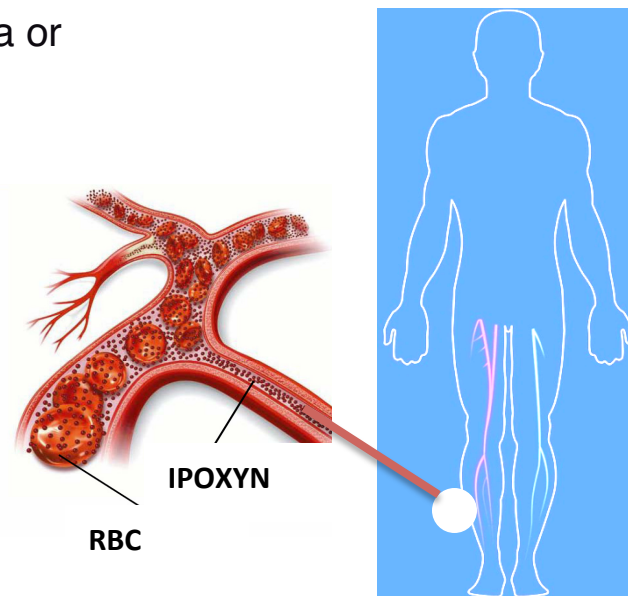


IPOXYN

- Carbohydrate-based intravenous solution that can potentially prevent necrosis, or cell death
- Treats ischemia, or lack of oxygen supply to living cells
- New chemical entity, not a biologic agent therefore strong regulatory position compared to biologic competitors
- Prevent amputation associated with lower limb ischemia or diabetic foot
- Contains oxygen rechargeable iron which picks up oxygen in the lungs
- 5,000 times smaller than red blood cell (RBC)
- Requires no blood type matching

Necrosis: localized death of living tissue.

Ischemia: deficient supply of blood to a body part, **leading to necrosis**



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IPOXYN: Market Opportunity

Facts and Figures

Global Market: \$30 billion*

Indications in which **necrosis** occurs:

- Stroke
- Heart Disease
- Trauma
- Anemia
- Stroke is a leading cause of death in the US
- Over 800,000 people die in the US each year from cardiovascular disease and strokes
- Kidney Failure
- Diabetic Foot
- Surgery

Competitive Advantage

- No current drug available to treat or prevent necrosis
- Ischemia currently treated by high pressure (hyperbaric) chamber
- All oxygen therapeutic drugs have failed in FDA trials
- IPOXYN is stable and does not scavenge Nitric Oxide
- Stable at room temperature

Source: Center for Disease Control and Prevention; *bcc Research

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2012 Milestones Achieved

- ✓ PAZ320 Phase II clinical trial results indicated no serious adverse events
- ✓ PAZ320 Phase II clinical trial results show 40% reduction in the elevation of post-meal blood sugar

Product	2013	2014	2015	2016
PAZ320	<ul style="list-style-type: none"> • Publish Phase II Data • Initiate Phase III international trial • Initiate Phase II trial in France • Submitted Pre-IND meeting request to FDA for PAZ320 	<ul style="list-style-type: none"> • Pivotal study initiation • Phase III study finalized 	<ul style="list-style-type: none"> • Clinical studies report finalized 	<ul style="list-style-type: none"> • New Drug Application (NDA)
IPOXYN	<ul style="list-style-type: none"> • Initiate pre-clinical experiments 	<ul style="list-style-type: none"> • Short term toxicity studies • Pre-IND meeting with FDA 	<ul style="list-style-type: none"> • IND application 	<ul style="list-style-type: none"> • First in human study indication

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Highlights

- Mission to become a leading pharmaceutical company focused on the development and commercialization of novel products for treatment of diabetes and inflammatory diseases
- Uniquely situated to take advantage of the increasing demand for innovative carbohydrate drug design
- IP portfolio places the Company ahead of the curve in carbohydrate technology development
- Team with extensive expertise in regulatory and clinical development, with multiple submissions and approvals to the FDA