



39TH ANNUAL J.P. MORGAN HEALTHCARE CONFERENCE

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Nasdaq: GWPH

LILLY
Living with Dravet syndrome

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Forward-Looking Statements

This presentation contains forward-looking statements that are based on our management's beliefs and assumptions and on information currently available to management. Forward-looking statements include information about our current expectations for future events, including potential results of operations, the timing of clinical trials, the timing of regulatory filings and approvals, the timing and outcomes of regulatory or intellectual property decisions, demand for our commercially available products and products in development and the clinical benefits, safety profile and commercial potential and potential pricing of Sativex[®], Epidiolex[®], and any product candidates. These forward-looking statements are subject to known and unknown risks, uncertainties, assumptions, including those associated with COVID-19 pandemic, and other factors that could cause our actual results, performance or achievements to be materially different than any future results, performance or achievements expressed or implied by the forward-looking statements. Forward-looking statements represent our management's beliefs and assumptions only as of the date of this presentation. You should read our most recent filings with the Securities and Exchange Commission including our Annual Report on Form 10-K and our Quarterly Reports on Form 10-Q, including the Risk Factors set forth therein and the exhibits thereto, and our subsequent filings with the Securities and Exchange Commission, completely and with the understanding that our actual future results may be materially different from what we expect. Except as required by law, we assume no obligation to update these forward-looking statements publicly, or to update the reasons actual results could differ materially from those anticipated in the forward-looking statements, even if new information becomes available in the future.



THE GLOBAL LEADER IN CANNABINOIDS

20+ Years Expertise
Proven Track Record

GW
pharmaceuticals


2020 Key Achievements

PROGRESSED



Progressed an active R&D program and robust pipeline

PREPARED




Prepared Nabiximols for US development and submission

EXPANDED



Expanded indication and achieved commercial success of Epidiolex[®]

GREW



Grew global reach and established EU market presence

TAKING CANNABINOID LEADERSHIP TO THE NEXT LEVEL

EPIDIOLEX



Highly-purified botanicals

NABIXIMOLS



Complex botanicals

NOVEL PRODUCT CANDIDATES



Next-generation molecules

PLANT DERIVED

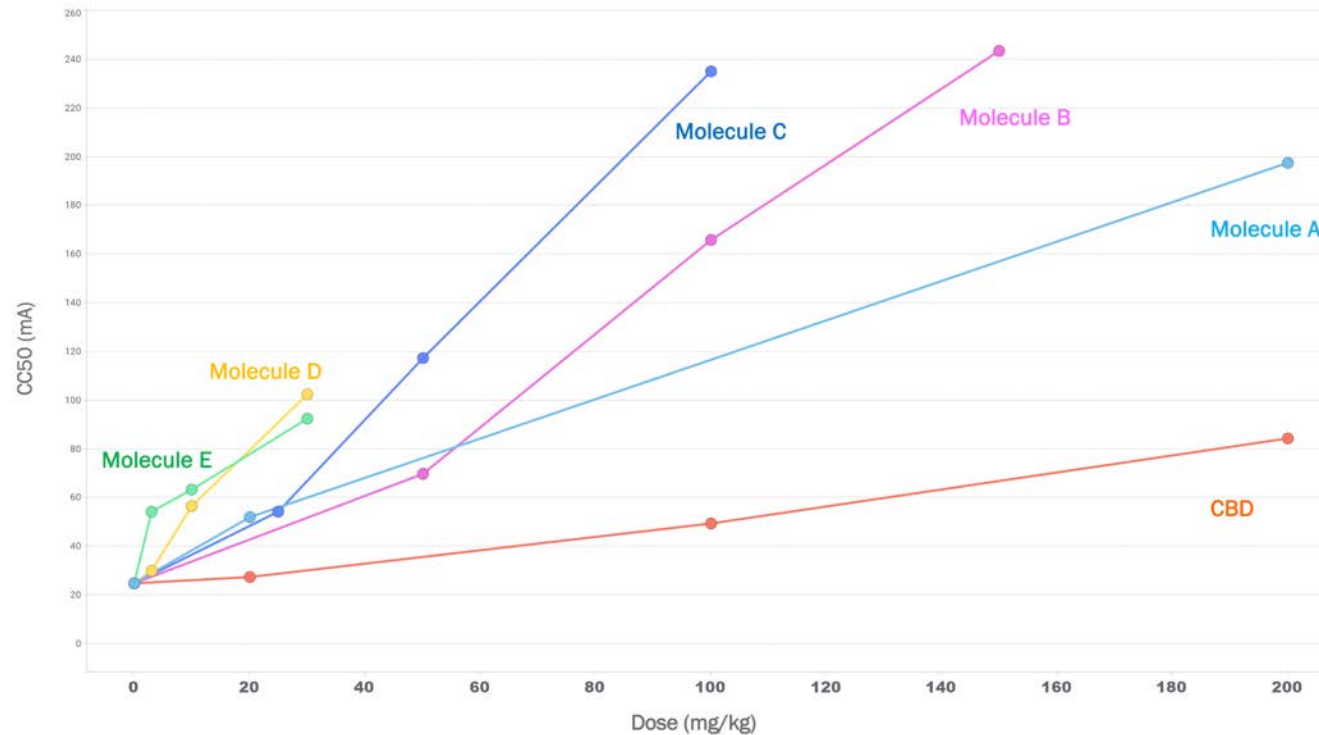
BEYOND THE PLANT

As the leader in cannabinoid science, GW has the greatest scientific understanding of pharmacological effects, unique formulation and drug delivery requirements and related exclusivity of cannabinoids.

Enhanced pharmaceutical properties
Increased potency
Composition of matter IP protection
Preserving cannabinoid efficacy and safety characteristics

First Novel Product Candidate Ready to Enter the Clinic in 2021

Novel product candidates demonstrate improved pharmaceutical and pharmacodynamic (efficacy) properties in relevant seizure model



Multiple new molecules already identified with enhanced pharmaceutical properties, signals of increased potency with patents filed.

Novel product candidates



Assess pharmacology



Physicochemical properties, including solubility



Metabolic assessment



Improved efficacy and potency in model of acute generalized seizures

Epidiolex[®] (cannabidiol) Oral Solution

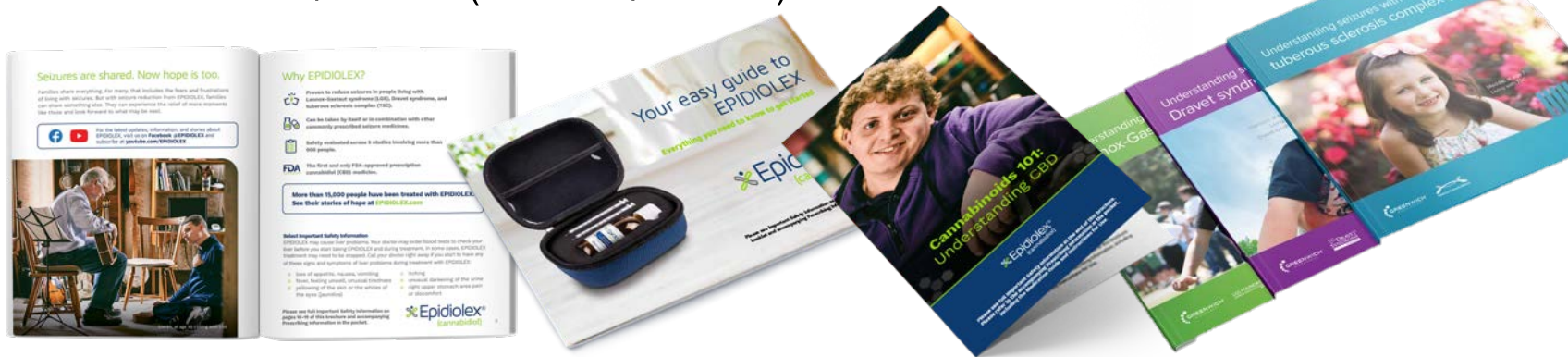
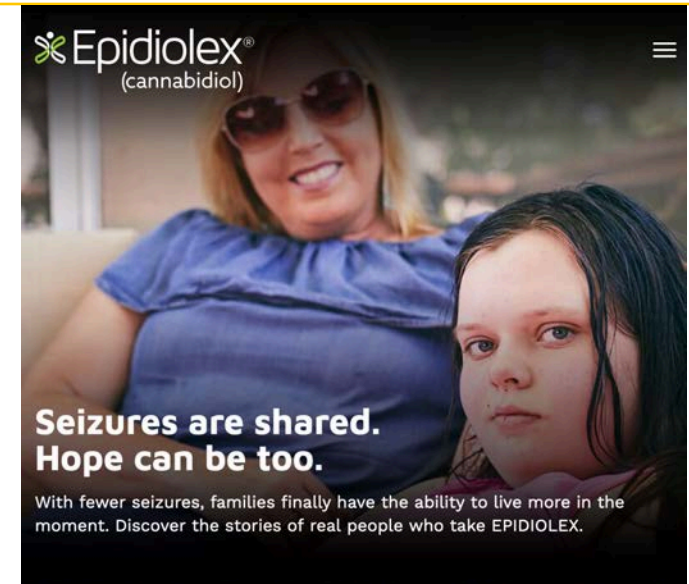
PIPER

Living with Dravet syndrome

Continued US Commercial Success and Growth



- First FDA-approved plant-derived cannabinoid; new MOA
- Descheduled by DEA
- Indicated to treat seizures associated with LGS, Dravet syndrome and TSC
 - Broad spectrum efficacy demonstrated in 3 approved indications
- Indicated in patients 1 year of age and older
- High level of awareness and intent to prescribe
- Unaudited 2020 Epidiolex global net product sales increased over 70% to \$510m (2019: \$296m)



Tuberous Sclerosis Complex (TSC) Launch and Opportunity

- Approved by FDA in Q3 2020
- TSC represents approx. 30,000 additional indicated patients in US
 - Epilepsy is present in >90% of patients with TSC
 - TSC affects 40-50K in the US
 - >60% of individuals with seizures associated with TSC do not achieve seizure control with standard treatments
 - Leading cause of genetic epilepsy, often occurring in first year of life
- Primary seizure types in TSC are focal and generalized
- EMA submission accepted for review



EPIDIOLEX is now approved for the treatment of seizures associated with TSC

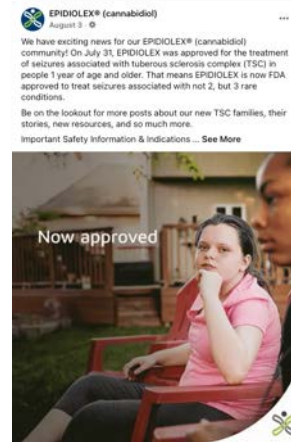
For people 1 year of age and older.

Dear Brent,

There's some big news in our community! EPIDIOLEX, the first and only prescription cannabidiol (CBD) to be approved by the FDA, was recently approved for the treatment of seizures associated with tuberous sclerosis complex (TSC). This means that EPIDIOLEX will be able to reach even beyond Lennox-Gastaut syndrome and Dravet syndrome—this means more hope.

EPIDIOLEX (cannabidiol)?
a prescription medicine that is used to treat seizures associated with Lennox-Gastaut syndrome, Dravet syndrome, or tuberous sclerosis complex in patients 1 year of age and older.

Important information I should know about EPIDIOLEX?
If you are allergic to cannabidiol or any of the ingredients in EPIDIOLEX may cause liver problems. Your doctor may order blood tests to check your liver before you start taking EPIDIOLEX and during treatment. Please see additional **Important Safety Information** below.



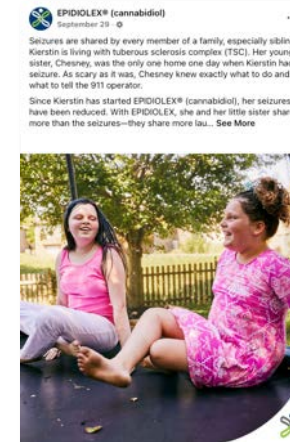
EPIDIOLEX® (cannabidiol)
August 3

We have exciting news for our EPIDIOLEX® (cannabidiol) community! On July 31, EPIDIOLEX was approved for the treatment of seizures associated with tuberous sclerosis complex (TSC) in people 1 year of age and older. That means EPIDIOLEX is now FDA approved to treat seizures associated with not 2, but 3 rare conditions.

Be on the lookout for more posts about our new TSC families, their stories, new resources, and so much more.

Important Safety Information & Indications... See More

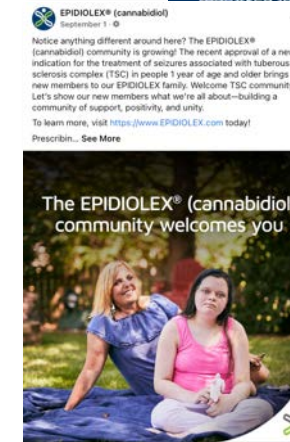
Now approved



EPIDIOLEX® (cannabidiol)
September 20

Seizures are shared by every member of a family, especially siblings. Kierstin is living with tuberous sclerosis complex (TSC). Her younger sister, Chesney, was the only one home one day when Kierstin had a seizure. As scary as it was, Chesney knew exactly what to do and what to tell the 911 operator.

Since Kierstin has started EPIDIOLEX® (cannabidiol), her seizures have been reduced. With EPIDIOLEX, she and her little sister share more than the seizures—they share more life... See More



EPIDIOLEX® (cannabidiol)
September 1

Notice anything different around here? The EPIDIOLEX® (cannabidiol) community is growing! The recent approval of a new indication for the treatment of seizures associated with tuberous sclerosis complex (TSC) in people 1 year of age and older brings new members to our EPIDIOLEX family. Welcome TSC community! Let's show our new members what we're all about—building a community of support, positivity, and unity.

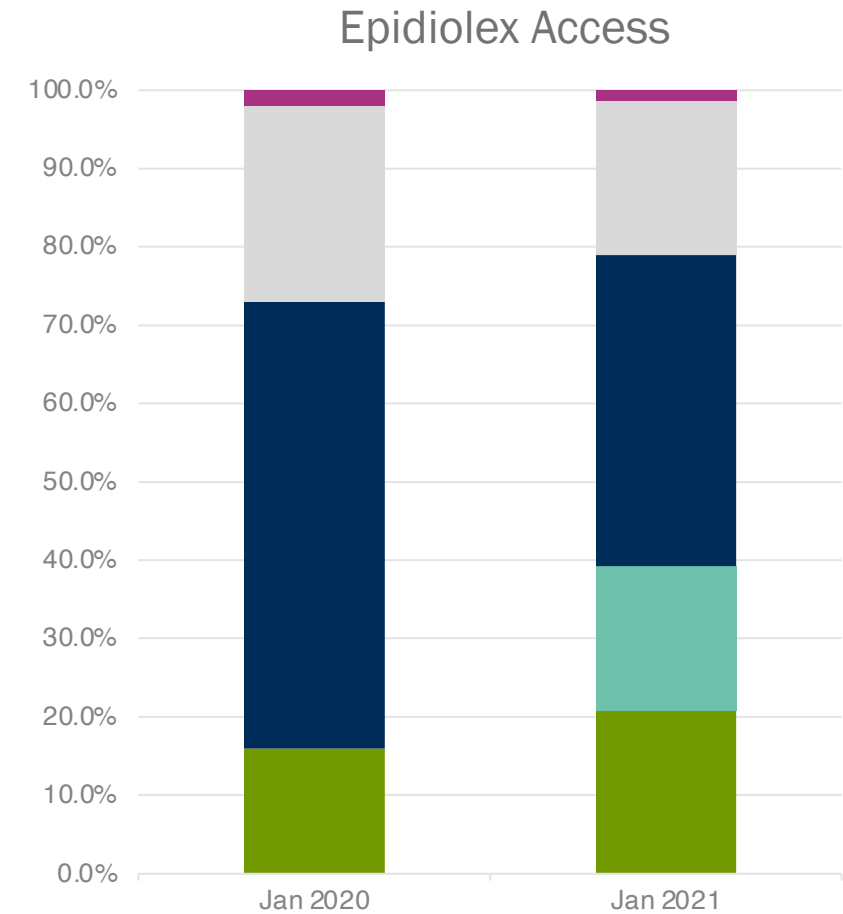
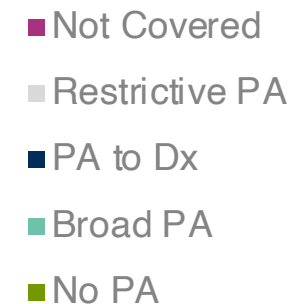
To learn more, visit <https://www.EPIDIOLEX.com> today!

Prescribin... See More

The EPIDIOLEX® (cannabidiol) community welcomes you

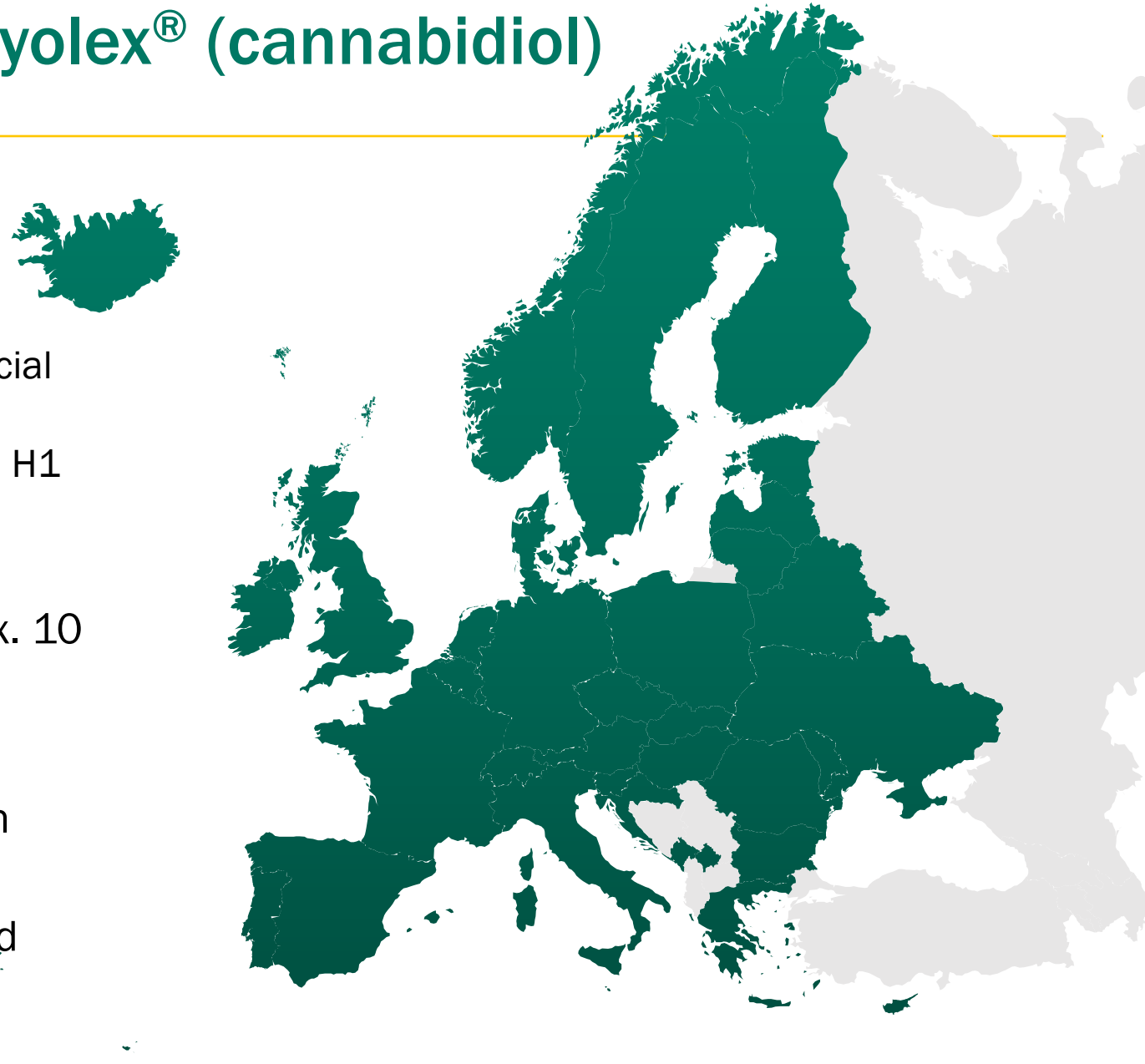
U.S. 2020 Payer Success

- Expanded payer coverage includes 85 million lives with no/broad prior authorization
- In 2020, increased access with payers having no/broad PA from 16% to 39% of US covered lives
- Approximately 2 out of 3 US Medicaid patients have broad access to Epidiolex
 - Medicaid is the largest payer for Epidiolex
- Commercial coverage expanding
 - Express Scripts - Epidiolex is a Preferred brand with lowest brand copay
 - Cigna
 - Others

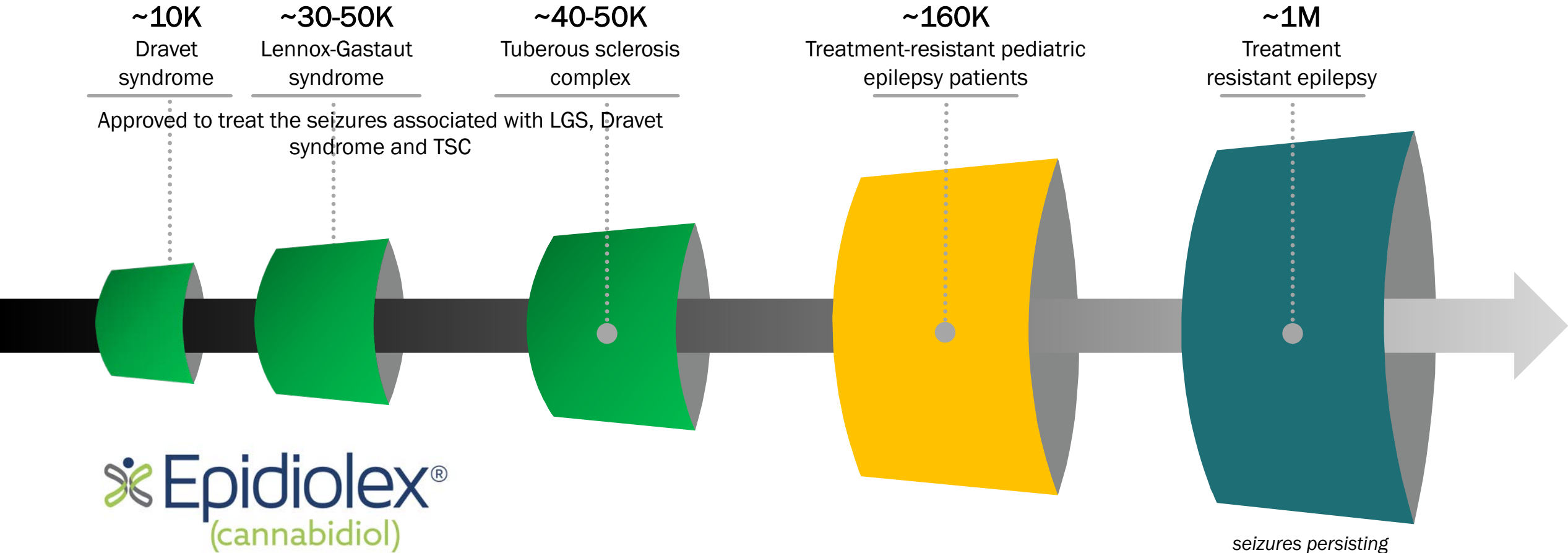


Expanded Global Reach of Epidyolex[®] (cannabidiol)

- EMA approval in September 2019
- First wave in 5 major markets
 - Germany - commercial launch in Q4 2019
 - UK - positive NICE recommendation with commercial launch in Q1 2020
 - France and Italy – commercial launches expected H1 2021
 - Spain – H1 launch
- Pricing and reimbursement progressing in approx. 10 additional European countries
- EMA TSC approval expected H1 2021
- Approved for use by TGA in Australia in 2020 with reimbursement recommended
- Opened offices in Berlin, Milan, Madrid, Paris and Amsterdam



Significant Unmet Need Remains For Further Research



Camfield CS, et al. *Epilepsia*. 1996;37(1):19-23; US Department of Commerce. <https://www.census.gov/prod/3/98pubs/p23-194.pdf>. 1997. Accessed May 29, 2018.; Camfield P, Camfield C. *Epilepsia*. 2007;48(6):1128-1132.; Berg AT, et al. *Epilepsia*. 2000;41(10):1269-1275.; Wu YW, et al. *Pediatrics*. 2015;136(5):e1310-e1315.; Centers for Disease Control. <https://www.cdc.gov/mmwr/volumes/66/wr/mm6631a1.htm>. 2017. Accessed April 19, 2018.; Kwan P, Brodie MJ. *N Engl J Med*. 2000;342:314-319; Sander JW, *Epilepsia*. 1993;34(6):1007; Picot et al, 2008 ; Kwan P, Brodie MJ. *N Engl J Med*. 2000;342:314-319; Kwan P, Brodie MJ, *CNS Spectr*. 2004;9(2):110; Epilepsy Foundation, <https://pediatrics.aappublications.org/content/136/5/e1310>.

Epidiolex 2021: Building a Blockbuster



BUILD

Build on positive physician experiences to increase prescribing to appropriate patients



ACCELERATE

Accelerate adoption across a broader prescriber base



DRIVE

Continue expanded payer access momentum



IMPROVE

Continue to improve the patient Epidiolex experience



INCREASE

Increase penetration in the long-term care segment



EXECUTE

Execute on the TSC label expansion



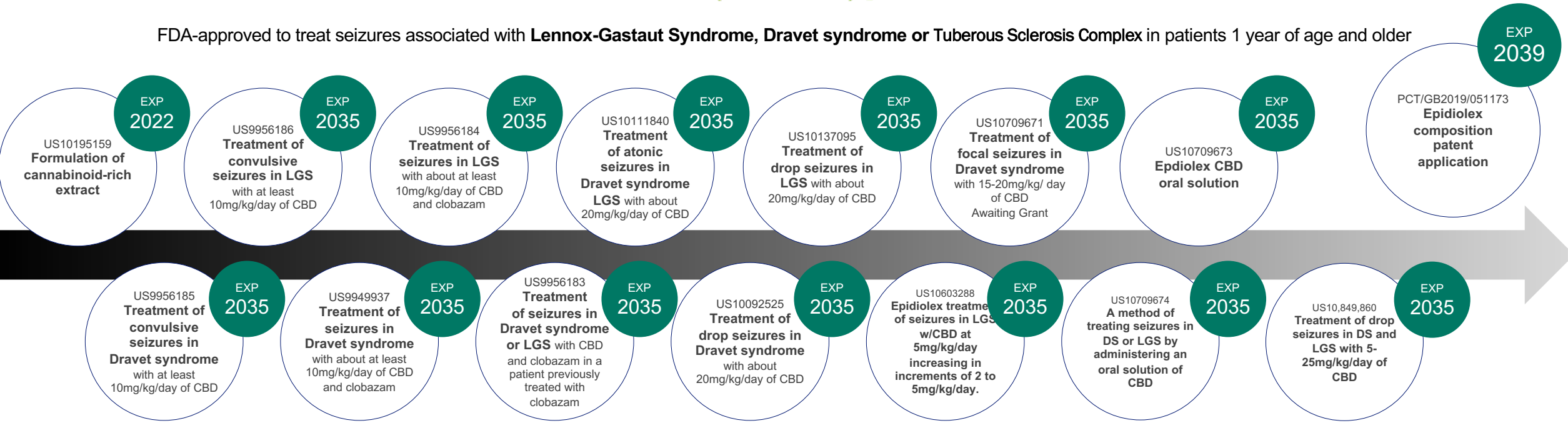
DEVELOP

Further indication development

Epidiolex® Patent Portfolio to 2035 Aligns with Indication



FDA-approved to treat seizures associated with **Lennox-Gastaut Syndrome, Dravet syndrome or Tuberous Sclerosis Complex** in patients 1 year of age and older



14 Orange Book-listed patents to date plus composition, TSC, and use patents in prosecution plus additional applications planned

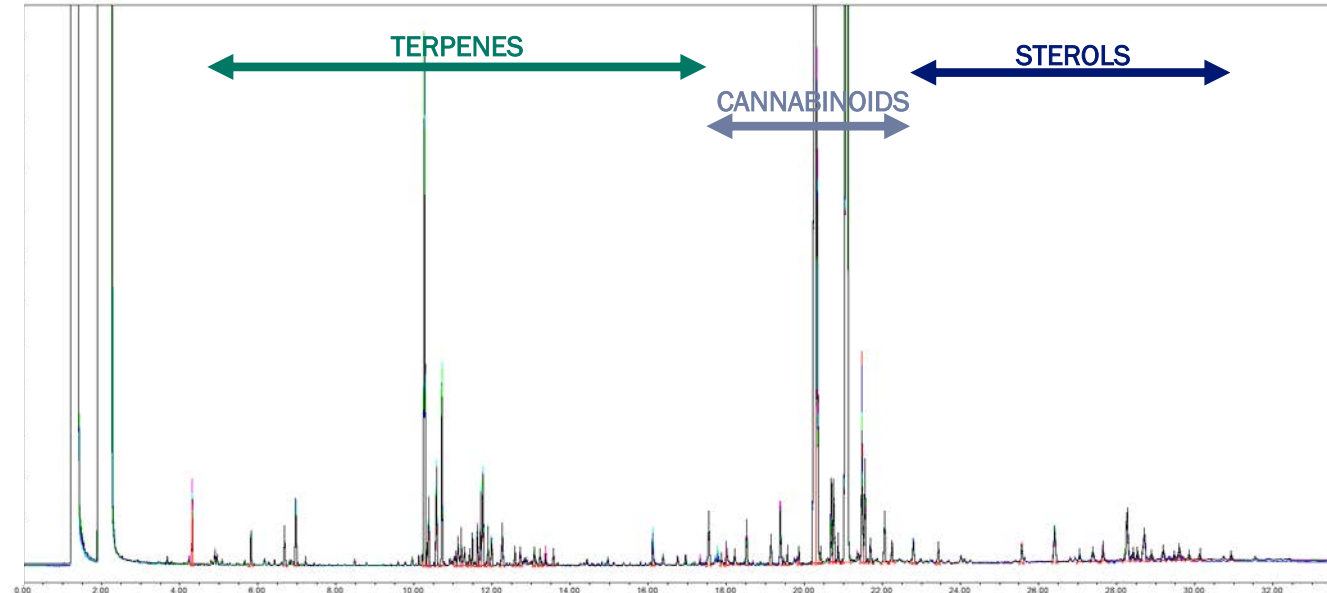


Nabiximols

Nabiximols: Next US Commercial Opportunity

- Derived from the whole cannabis plant containing a clinically proven, balanced dose of THC and CBD along with other cannabinoid and non-cannabinoid plant components
- Approved in >25 countries outside the US as Sativex® for the treatment of spasticity due to multiple sclerosis (MS); sold via marketing partners
- Near-term, reduced US risk opportunity in MS Spasticity
 - Positive efficacy, safety and abuse/diversion data
 - US pivotal clinical program now recruiting
- Broad potential in spasticity beyond MS
- Strong exclusivity due to complex botanical formulation
- US commercial rights owned by GW

Nabiximols – Reproducible Fingerprint, 5 Batches Overlaid



MS Spasticity Market (US)

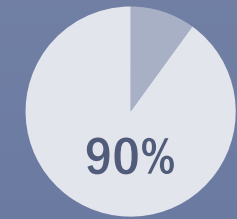
- Spasticity occurs in up to 84% of MS patients ¹
- Despite current treatment, 1/3 live with uncontrolled spasticity ²
- No new oral anti-spasticity medicines approved in over 20 years; MS disease modifying treatments have not shown evidence of helping to relieve symptoms
- Spasticity significantly impacts daily function – reduced mobility and inability to perform daily tasks, including walking and driving ³
- 26–50% of MS patients are self-medicating with cannabis ⁴

Sources:

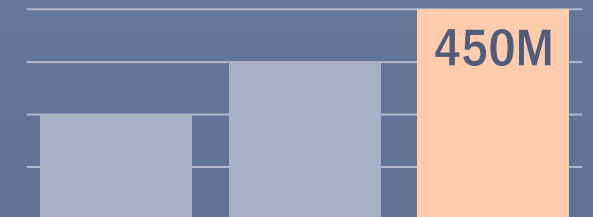
- 1.) Rizzo M, Hadjimichael O, Preiningerova J, Vollmer T. Prevalence and treatment of spasticity reported by multiple sclerosis patients. *Multiple Sclerosis Journal*. 2004;10(5):589-595.
- 2) Pozzilli C. Advances in the management of multiple sclerosis spasticity: experiences from recent studies and everyday clinical practice. *Expert review of neurotherapeutics*. 2013;13(12 Suppl):49-54. *Mult Scler*. 2004 Oct; 10(5):589-95.
- 3.) Bethoux F, Marrie RA. A cross-sectional study of the impact of spasticity on daily activities in multiple sclerosis. *The Patient-Patient-Centered Outcomes Research*. 2016;9(6):537-546.
- 4.) 2014 NARCOMS survey, Rizzo et al. *Mult Scler* 2004, Rizzo et al. *Mult Scler* 2004; Braley et al. *Mult Scler J Exp Transl Clin* 2020.
- 5.) Clearview MS Spasticity Landscape Assessment August 2019

NABIXIMOLS OPPORTUNITY IN MS SPASTICITY

Strong physician and patient
enthusiasm for product use ⁵



90% of physicians and patients
highly to moderately interested ⁵



**US market potential estimated
to be ~\$450 million**

Nabiximols: Demonstrated Efficacy in MS Spasticity

	STUDY 1, Colin <i>et al</i>	STUDY 2, Novotna <i>et al</i>	STUDY 3, Markova <i>et al</i>
STUDY DESIGN	6 Week, Double-blind, Randomized Parallel Group Study	Enriched Randomized Withdrawal Study Design	Enriched Study Design
SAMPLE SIZE	N = 189	Part A: N = 572 Part B: N= 241	Part A: N = 191 Part B: N= 106
PRIMARY ENDPOINTS	Primary: NRS-spasticity (mean change) Key secondary: NRS-spasticity (responder) Ashworth scale, spasms	Primary: NRS-spasticity (mean change) Key secondary: NRS-spasticity (responder) spasms, modified Ashworth scale (mAS)	Primary: NRS spasticity (responder) Key secondary: NRS-spasticity (mean change), modified Ashworth scale
PRIMARY ENDPOINT P-VALUE	P=0.048	P=0.0002	P<0.0001

Collin, et al. Eur J Neurol 2007; 14:290-96

Novotna, et al. Eur J Neurol 2011; 18(9):1122-31

Markova, et al. Int J Neurosci 2019; 129(2): 119-128

Agreed Path Forward With The FDA

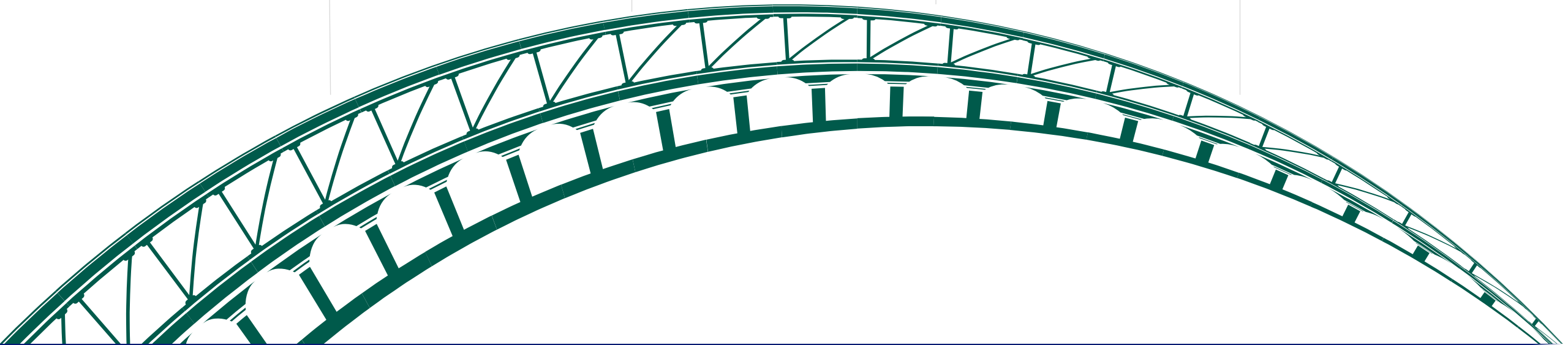
EUROPEAN EFFICACY TRIALS PRESENTED TO FDA

DATA HARMONIZED TO NDA REQUIREMENTS

BRIDGE FROM NRS TO PROXIMATE SPASTICITY ENDPOINT

EXECUTE MULTIPLE PIVOTAL TRIALS: FIVE SHOTS ON GOAL

DISCUSS EXPANDING INDICATION



Five Phase 3 Studies; NDA Filing on First Positive Readout

Sample Size	Design	Endpoint	Start	Target data readout
52	3-week treatment period, placebo-controlled cross-over	Muscle Tone (LLMT-6)	Dec 2020	2021
190	3-week treatment period, placebo-controlled cross-over	Muscle Tone (LLMT-6)	Q1 2021	2022
36	Pre-post dose in prescribed responders, placebo-controlled cross-over	Muscle Tone (LLMT-6)	Q1 2021	2021/2022
446	Double-blind, parallel, placebo controlled, 12-week treatment period	Spasm frequency	Oct 2020	2022
227 randomized	Part A: 4-week run-in to identify responders; Part B: 12-week double-blind, parallel, placebo-controlled randomized withdrawal	Spasm frequency	H1 2021	2022/3

Opportunity to Achieve Broad Spasticity Label As Well as Other Lifecycle Opportunities

- Broader spasticity population
 - >3M U.S. patients including spinal cord injury, post-stroke, ALS, traumatic brain injury, cerebral palsy
- Spasticity associated with Spinal Cord Injury (SCI)
 - Approx. 250K chronic SCI patients (~65%) suffer from spasticity
 - Likely single pivotal trial required
 - Sales potential ~\$350M
- Post Traumatic Stress Disorder (PTSD)
 - Impacting ~11.7M people with ~55% diagnosed
 - Anxiety is one of the top 3 reasons for self-medication with cannabis
 - Nabiximols offers potential to reduce sleep disturbance symptoms, anxiety and irritability

Sources: DiPiro. Spinal Cord. 2018; McGuire. Spasticity: Diagnosis and Management, 2011; Nicholson. Muscle Nerve. 2018; AANS Website; UpToDate; Physician Interviews; ClearView Analysis. Goldstein. Soc Psychiatry Psychiatr Epidemiol 2016; Kessler. Arch Gen Psychiatry. 2012; Kessler. Arch Gen Psychiatry. 2005; UpToDate; Physician Interviews; ClearView Analysis



Corporate Information

Q4 Results and Key Financial Data (Unaudited)

\$148M

Q4 2020
net revenue

\$486M

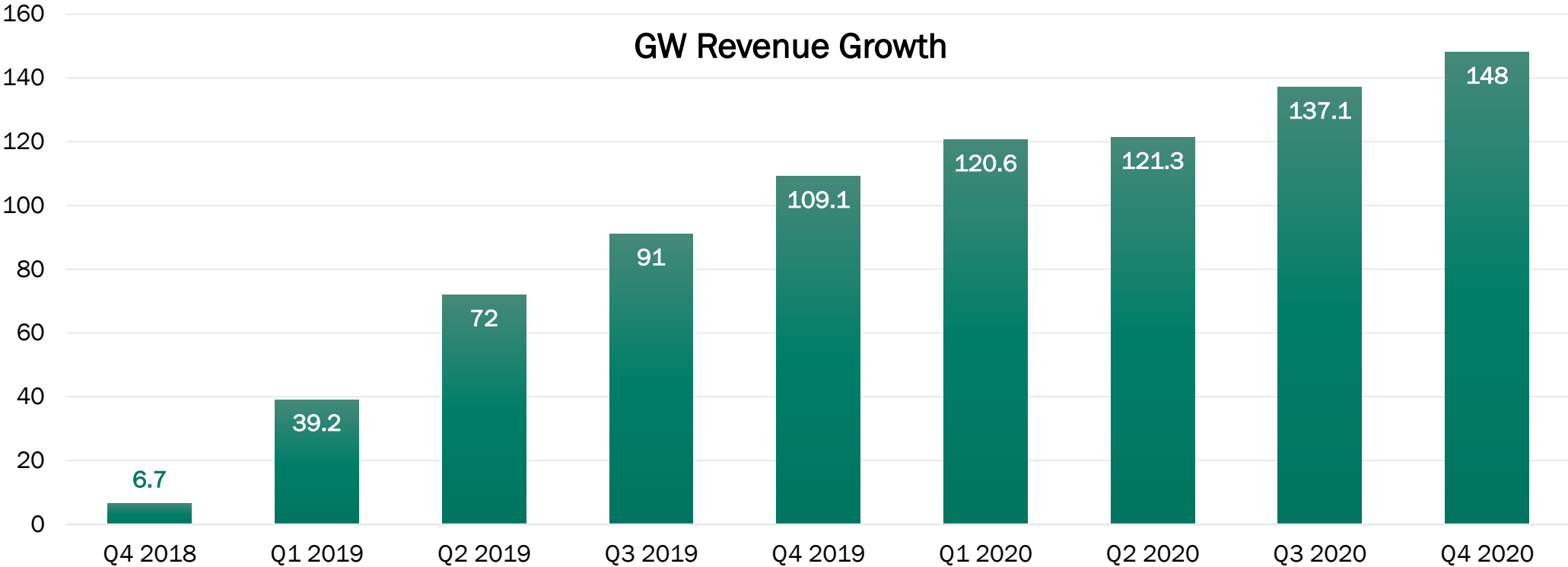
Cash at
December 31, 2020

\$530M–550M

FY 2020 R&D and
SG&A expense guidance

\$20M–25M

FY 2020 capital
expense guidance



2021 Key Priorities

GROW



Grow Commercial Success of Epidiolex®

PREPARE



Prepare Nabiximols for approval and commercial launch

ADVANCE



Advance R&D Program and Robust Pipeline

EXPAND



Expand Global Reach of products and pipeline



THANK YOU!

www.gwpharm.com



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Living with Dravet syndrome