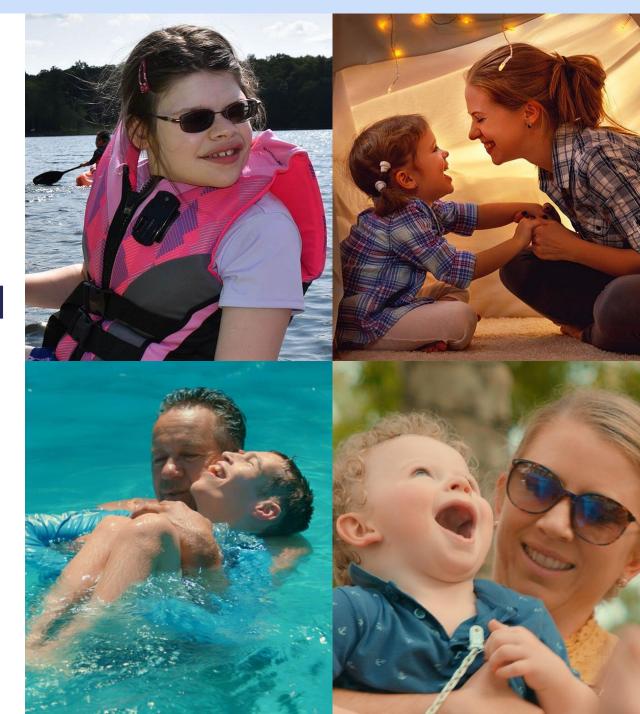


Neuren and Acadia Expand Global Partnership for trofinetide (DAYBUETM)

14 July 2023

IMPROVING THE LIVES OF PEOPLE WITH NEURODEVELOPMENTAL DISABILITIES



Forward looking statements

This presentation contains forward looking statements that involve risks and uncertainties. Although we believe that the expectations reflected in the forward looking statements are reasonable at this time, Neuren can give no assurance that these expectations will prove to be correct. Actual results could differ materially from those anticipated. Reasons may include risks associated with drug development and manufacture, risks inherent in the regulatory processes, delays in clinical trials, risks associated with patent protection, future capital needs or other general risks or factors.





Transaction to maximize value of trofinetide and NNZ-2591

Expands Acadia's exclusive trofinetide licence to worldwide, for up to additional US\$527m to Neuren (including US\$100m up-front) plus mid-teens to low-20s % royalties on net sales outside North America

Leverages Acadia's unique knowledge and expertise from successful DAYBUE development and commercialization in the US and the established supply chain; Acadia responsible for all costs

Unlocks NNZ-2591's potential application in Rett and Fragile X globally, with separate sales milestones and royalties to Neuren identical to trofinetide inside and outside North America

Significantly strengthens Neuren's position to explore all strategic options, with cash at 30 June 2023 plus the up-front payment¹ A\$226 million

¹ Assuming AUD/USD exchange rate of 0.68 and US withholding tax of 5% for the up-front payment



Transaction economics to Neuren for trofinetide outside North America

	US\$100m	upfront payment
	US\$35m	following 1st commercial sale in Europe
	US\$15m	following 1st commercial sale in Japan
	Up to US\$14m	in relation to 1st commercial sale of a 2 nd indication in Europe and Japan
Total payments unrelated to sales performance	US\$164m	
unrelated to sales		On achievement of escalating annual net sales thresholds

If Acadia sub-licenses any region within the first 2 years, Neuren is entitled to a share of any upfront and development milestones received by Acadia. Creditable against future payments to Neuren in the sub-licensed region.



Trofinetide North America Economics for Neuren unchanged

Sales Milestones

Rett Syndrome

√ US\$10m in 2022 following acceptance of N	NDA for review
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√ US\$40m following 1st commercial sale in the US

US\$33m one third share of Priority Review Voucher awarded to Acadia (assuming market value US\$100m)

Fragile X Syndrome

Up to US\$55m

in relation to development milestones and 1st commercial sale in the US

Aggregate of all indications

Tiered Royalty	Rates	(%	of	net
sales) ¹				

Annual Net Sales	Rates	Net Sales in one calendar year	US\$m
≤US\$250m	10%	≥US\$250m	50
>US\$250m, ≤US\$500m	12%	≥US\$500m	50
>US\$500m, ≤US\$750m	14%	≥US\$750m	100
>US\$750m	15%	≥US\$1bn	150
		Total	350

Example calcumilestones	Example calculations of royalty/sales milestones		
Annual Net	Annual	Total Sales	

Sales in NA	Royalty	Milestones Earned
US\$500m	US\$55m	US\$100m
US\$750m	US\$90m	US\$200m
US\$1bn	US\$128m	US\$350m

¹ Royalty rates payable on the portion of annual net sales that fall within the applicable range



Value of NNZ-2591 enhanced

Before Transaction Post Transaction Exclusive worldwide licence to Acadia for Rett and Fragile X syndromes only Rett Rett **Partnership** Global with Acadia . Potential future payments to Neuren for NNZrestrictions 2591 in Rett and Fragile X syndromes Fragile X Fragile X identical to the payments for trofinetide inside and outside North America Phelan-Phelan-**McDermid McDermid** Pitt Hopkins Pitt Hopkins Neuren retains worldwide rights to NNZ-2591 in all other indications Angelman Angelman Ongoing Phase 2 clinical trials in Phelan-Neuren Restrictions in McDermid, Pitt Hopkins, Angelman and North America Prader-Willi syndromes Prader-Willi Prader-Willi only if Acadia First top-line results expected in Dec 2023 develops trofinetide for I Other Other an indication



Transforming catalysts in 2023

Commercial





Trofinetide NA

Trofinetide RoW

NNZ-2591

Development

- ✓ DAYBUE for Rett syndrome approved by FDA
- ✓ Priority Review Voucher awarded to Acadia
- ✓ First US commercial sale US\$40m milestone payment
- Quarterly royalties on net sales
- Priority Review Voucher value one third share estimated as US\$33m
- ✓ Global trofinetide partnership with Acadia
- Receive US\$100m upfront payment from Acadia
- ✓ Initiate Prader-Willi syndrome Phase 2 trial
- ✓ Complete enrolment in Phelan-McDermid syndrome Phase 2 trials
- Complete enrolment in Pitt Hopkins and Angelman syndrome Phase 2 trials
- Top-line results for Phelan-McDermid syndrome in Dec 2023



