

Bell Potter Healthcare Conference Presentation

Melbourne, Australia; 18 November 2024: Starpharma (ASX: SPL, US OTC: SPHRY), an innovative biotechnology company with two decades of experience in advancing dendrimer technology from the lab to the patient, provides a copy of the presentation that will be delivered by CEO Cheryl Maley at the virtual Bell Potter Healthcare Conference today.

About Starpharma

Starpharma ASX: SPL, US OTC: SPHRY) is an innovative biotechnology company with two decades of experience in advancing dendrimer technology from the lab to the patient. Our mission is to help patients with significant illnesses, such as cancer, achieve improved health outcomes and quality of life through the application of our unique dendrimer technology.

Dendrimers are precise, synthetically manufactured, nanoscale molecules. Their unique properties—including their size, structure, high degree of branching, polyvalency, and water solubility—are advantageous in medical and pharmaceutical applications.

Starpharma's portfolio of dendrimer-based products includes three clinical-stage DEP® (dendrimer enhanced product) assets, preclinical radiopharmaceutical assets, research collaborations, and three commercially marketed over-the-counter (OTC) products.

For more information about Starpharma, visit www.starpharma.com or connect with Starpharma on [LinkedIn](#).

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Disclosure

This ASX Announcement was authorised for release by the Chair, Mr Rob Thomas.

Forward-Looking Statements

This document contains certain forward-looking statements, relating to Starpharma's business, which can be identified by the use of forward-looking terminology such as "promising", "plans", "anticipated", "will", "project", "believe", "forecast", "expected", "estimated", "targeting", "aiming", "set to", "potential", "seeking to", "goal", "could provide", "intends", "is being developed", "could be", "on track", or similar expressions, or by express or implied discussions regarding potential filings or marketing approvals, or potential future sales of product candidates. Such forward-looking statements involve known and unknown risks, uncertainties and other factors that may cause actual results to be materially different from any future results, performance or achievements expressed or implied by such statements. There can be no assurance that any existing or future regulatory filings will satisfy the FDA's and other authorities' requirements regarding any one or more product candidates, nor can there be any assurance that such product candidates will be approved by any authorities for sale in any market or that they will reach any particular level of sales. In particular, management's expectations regarding the approval and commercialization of the product candidates could be affected by, among other things, unexpected trial results, including additional analysis of existing data, and new data; unexpected regulatory actions or delays, or government regulation generally; our ability to obtain or maintain patent or other proprietary intellectual property protection; competition in general; government, industry, and general public pricing pressures; and additional factors that involve significant risks and uncertainties about our products, product candidates, financial results and business prospects. Should one or more of these risks or uncertainties materialize, or should underlying assumptions prove incorrect, actual results may vary materially from those described herein as anticipated, believed, estimated, or expected. Starpharma is providing this information as of the date of this document and does not assume any obligation to update any forward-looking statements contained in this document as a result of new information, future events or developments or otherwise. Clinical case studies and other clinical information given in this document are given for illustrative purposes only and are not necessarily a guide to product performance and no representation or warranty is made by any person as to the likelihood of achievement or reasonableness of future results. Nothing contained in this document, nor any information made available to you is, or shall be relied upon as, a promise, representation, warranty or guarantee as to the past, present or the future performance of any Starpharma product.



ASX : SPL
US OTC : SPHRY

Delivering Meaningful Patient Outcomes with Advanced Dendrimer Technology

Bell Potter Healthcare Conference | 18 November 2024

CEO, Cheryl Maley

Disclaimer and 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, Starpharma 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.

Our Mission

“To help patients with significant illnesses, such as cancer, achieve improved health outcomes and quality of life through the application of our unique dendrimer technology.”

Starpharma is an Innovative Australian Biotechnology Company with 20 Years of Experience in Advancing Dendrimer Technology from the Lab to the Patient

Ticker Symbol	ASX: SPL US OTC: SPHRY
Industry/Sector	Healthcare, Pharmaceuticals, Biotechnology
Market Capitalisation	~A\$40M
Share Price	~A\$0.10
Total Ordinary Shares On Issue	417M
Cash Balance (30 Sept 2024)	A\$24.0M

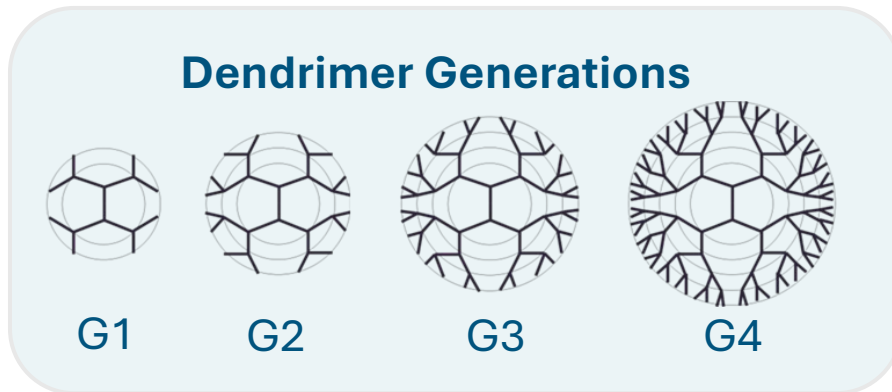


Starpharma's DEP[®] Platform Technology: What Is It and How Does It Benefit Patients?



Starpharma – Founders and Experts in Dendrimer Drug Delivery

Dendrimers are highly branched (tree-like) macromolecules with a well-defined, 3D structure



- Concentric layers of lysine monomers
- Drugs, payloads, and/or targeting moieties attached via tailored linker strategies to achieve enhanced tumour targeting and pharmacokinetics (PK)
- Easily scalable, precisely manufactured, and Good Manufacturing Practice (GMP) certified



Clinically validated technology

More than 350 patients treated with DEP® across multiple clinical programs.



Strong intellectual property position

19 active patent families with over 150 granted patents and more than 40 patent applications pending.



Uniquely experienced team

Expertise in dendrimer science.
Staff of ~40 people.

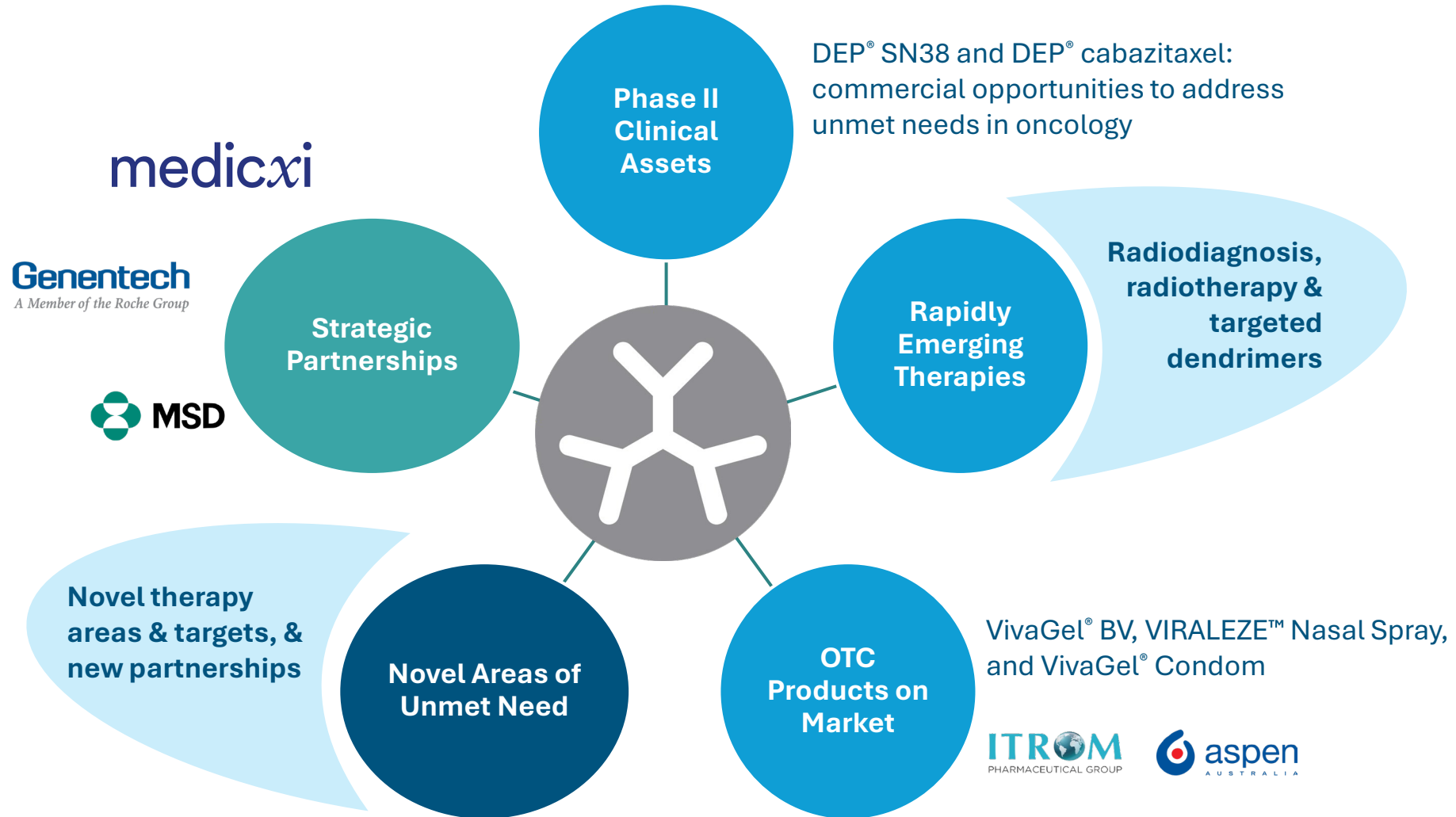


Pipeline of products and partnerships

Portfolio includes clinical-stage assets, early-stage research, partnerships, and commercial products.

Starpharma's DEP[®] Platform Technology: Versatile and Multifunctional for Delivery of Therapeutics and Diagnostics

Multiple Revenue Streams: Maximising Shareholder Returns



Starpharma's Oncology Portfolio: Addressing Areas of High Unmet Clinical Need

Three Clinical Assets and Three Pipeline Assets

Product	Target indication	Research	Preclinical	Phase I	Phase II	Phase III	Strategy
DEP [®] SN38	Ovarian and colorectal	Phase II results reported					License/co-develop – ovarian, colorectal
DEP [®] cabazitaxel	Prostate and ovarian	Phase II results reported					License – prostate, ovarian
DEP [®] HER2 radiodiagnostic	Diagnostic						Optimise and accelerate to clinical
DEP [®] HER2 radiotherapeutic	Solid cancers						Advance to clinical
DEP [®] HER2 ADC	Solid cancers						Advance to preclinical
DEP [®] docetaxel	Pancreatic and other cancers	Phase II results reported					Lower priority

DEP[®] SN38 & DEP[®] cabazitaxel

Global Commercial Opportunity



DEP[®] SN38
DEP[®] cabazitaxel

Both assets were developed using the DEP[®] technology to improve existing oncology products.



Partnering

Starpharma has created value through proof-of-concept and is seeking to license both products.



Promising Phase II Results

Phase II studies for each asset showed promising results of improved tolerability over the original compounds and comparable or improved efficacy. Trials have generated promising anti-cancer efficacy in very late-stage patients who have been heavily pre-treated.



Value Proposition

For a partner, both assets provide opportunity for new indications, new markets, and product life cycle extension.

DEP[®] SN38 Phase II Results

Clinically meaningful outcomes were achieved for patients who were heavily pre-treated prior to entering the trial and had few options.

Promising efficacy in patients with irinotecan-treated CRC and platinum-resistant/refractory ovarian cancer.

Well-tolerated with mostly mild/moderate gastrointestinal AEs, no cholinergic toxicity.

DEP[®] cabazitaxel Phase II Results

Clinical benefit even in patients previously exposed to taxanes, including standard cabazitaxel.

Promising efficacy in patients with mCRPC, ovarian and gastro-oesophageal cancers.

Well-tolerated with mostly mild/moderate AEs, no routine steroid premedication.

Indication Evaluation



Advanced colorectal cancer



Platinum-resistant ovarian cancer



Metastatic castration-resistant prostate cancer



Platinum-resistant ovarian cancer



Advanced gastro-oesophageal cancer

DEP[®] SN38 Phase I/II Trial Patients – Advanced, Heavily Pre-treated, and Most CRC Patients had Progressed Following Prior Treatment with Irinotecan

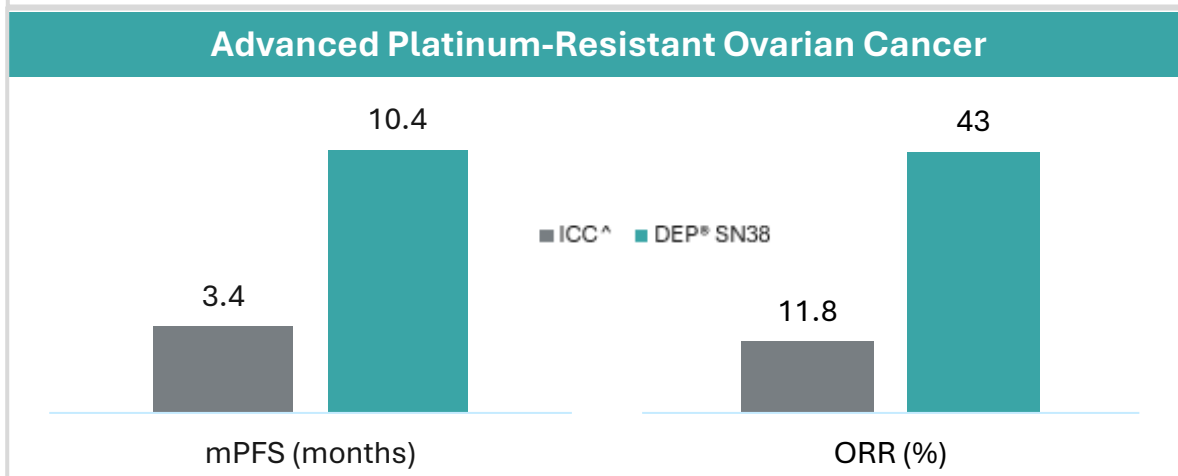
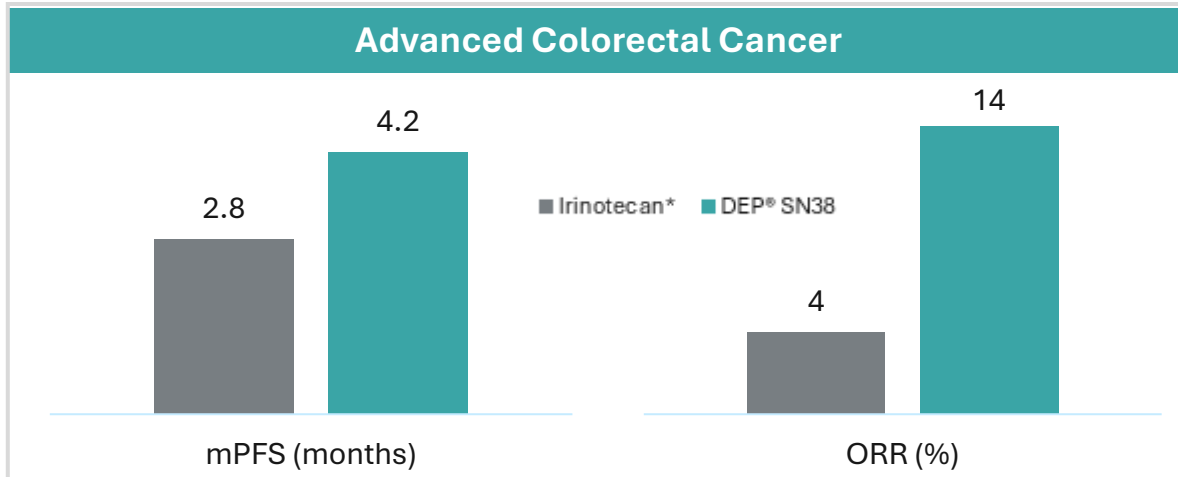
BASELINE CHARACTERISTICS		COLORECTAL	OVARIAN	PANCREATIC	BREAST	OTHER ¹	TOTAL
Subjects enrolled (n, %)		55 (48%)	23 (20%)	15 (13%)	8 (7%)	13 (11%)	114 (100%)
Subjects ongoing (n, %)		0 (0%)	1 (4%)	0 (0%)	0 (0%)	0 (0%)	1 (1%)
Age (years)	Median (range)	59 (31-78)	64 (42-74)	65 (48-76)	53 (42-66)	60 (38-73)	61 (31-78)
Sex (n, %)	Male	24 (44%)	0	8 (53%)	0	9 (69%)	41 (36%)
	Female	31 (56%)	23 (100%)	7 (47%)	8 (100%)	4 (31%)	73 (64%)
ECOG PS	0	23 (42%)	6 (26%)	6 (40%)	2 (25%)	-	40 (35%)
	1	32 (58%)	17 (74%)	9 (60%)	6 (75%)	2	74 (65%)
Stage at diagnosis	III	2 (4%)	4 (17%)	0 (0%)	0 (0%)	2 (15%)	8 (7%)
	IV	53 (96%)	19 (83%)	15 (100%)	8 (100%)	11 (85%)	106 (93%)
Prior systemic therapy (n, %)	Irinotecan	54 (98%)	0 (0%)	11 (73%)	0 (0%)	3 (23%)	68 (60%)
	Platinum	29 (53%)	23 (100%)	9 (60%)	0 (0%)	12 (92%)	73 (64%)
	Taxanes	0 (0%)	23 (100%)	2 (13%)	7 (88%)	9 (69%)	41 (36%)
Prior lines of therapy	Median (range)	4 (2-9)	6 (3 to 9)	2 (2 to 5)	7 (3 to 12)	3 (1 to 6)	4 (1 to 12)

¹Other cancer types included lung, upper gastrointestinal, and kidney.

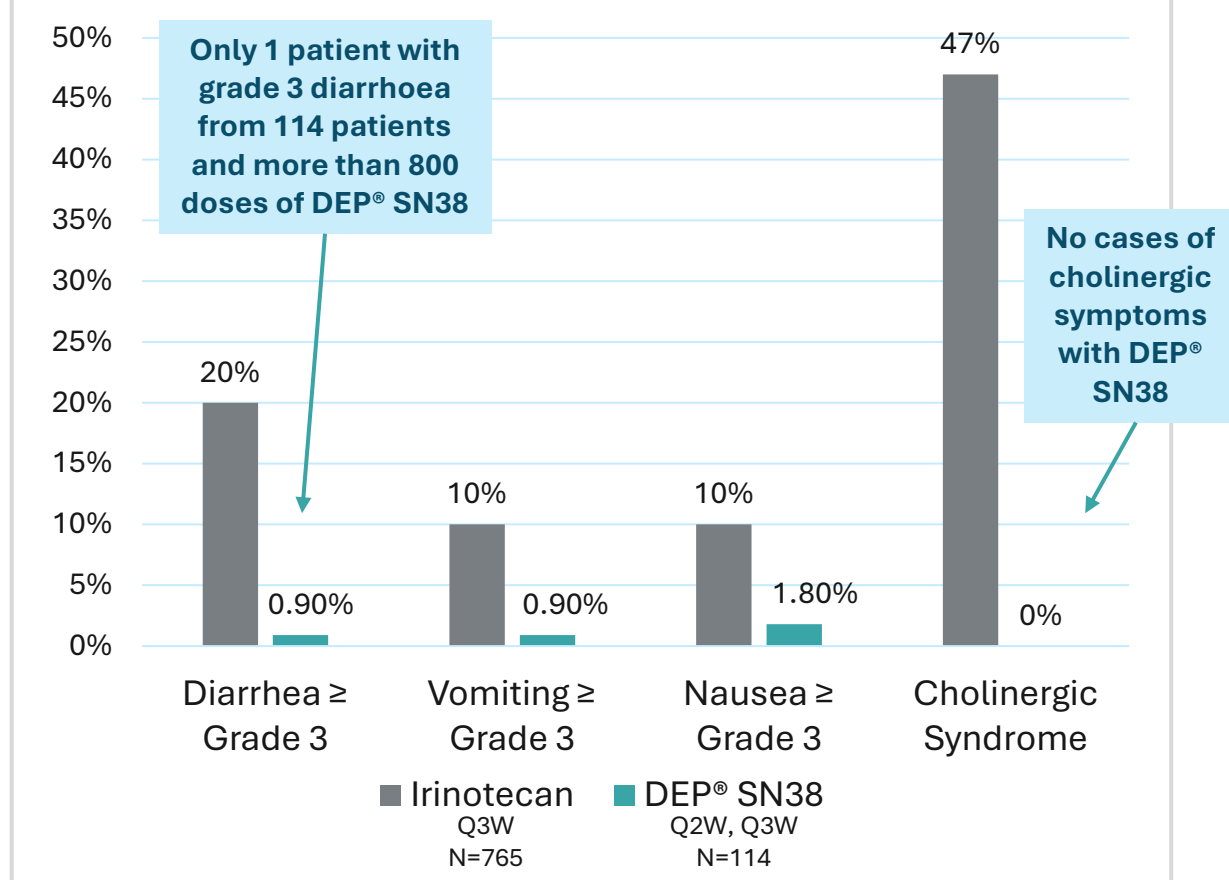
DEP[®] SN38 Phase II Study Shows Favourable Efficacy and Tolerability Data in Late-Stage Patients

Results Presented at the 2024 ASCO Annual Meeting

2024 ASCO[®]
ANNUAL MEETING



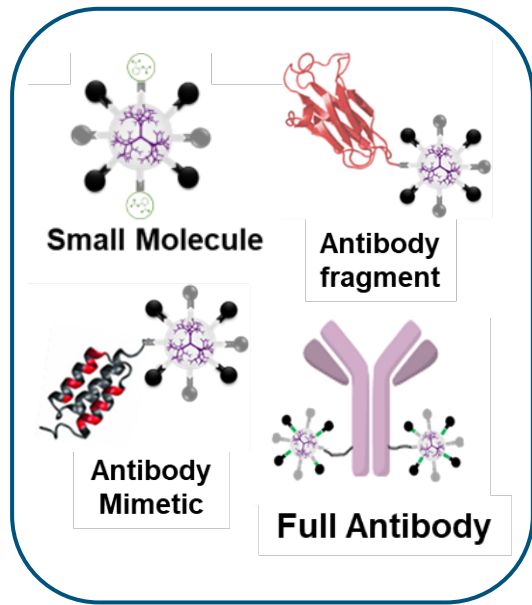
Gastrointestinal Toxicity Profile Significantly Improved with DEP[®] SN38 Treatment, Compared to Published Data on Irinotecan[#]



Data for DEP[®] SN38 in combination with 5-FU/LV; Full Phase II results reported in ASX Announcement dated 27 May 2024; *From published data on irinotecan in combination with 5-FU/LV, Tournigand et al., *Clin Oncol*, 2023, 41(19):3469-3477; # <https://www.medicines.org.uk/emc/product/6506>- UK SmPC April 2022

[#]From published data on ICC (investigator chemotherapy of choice) (pegylated liposomal doxorubicin, 11 paclitaxel, or topotecan), Pujade-Lauraine E, et al., *J Clin Oncol*, 2014, 32(13):1302-1308;

Benefits of Starpharma's DEP® Platform Technology Extend to Radiopharmaceuticals



Broad Applicability in Drug Development

Ability to use a wide range of targeting moieties

Site-specific attachment of dendrimer on targeting moiety

DEP® dendrimers are precisely manufactured and easily scalable

Drug-linker strategy flexibility

Flexibility in chelator type

Can select drug payload and radioisotope for the desired application



Key Characteristics Valued by Collaborators

Addressing the Biodistribution Challenges of Current Approaches with DEP® Radiotheranostics

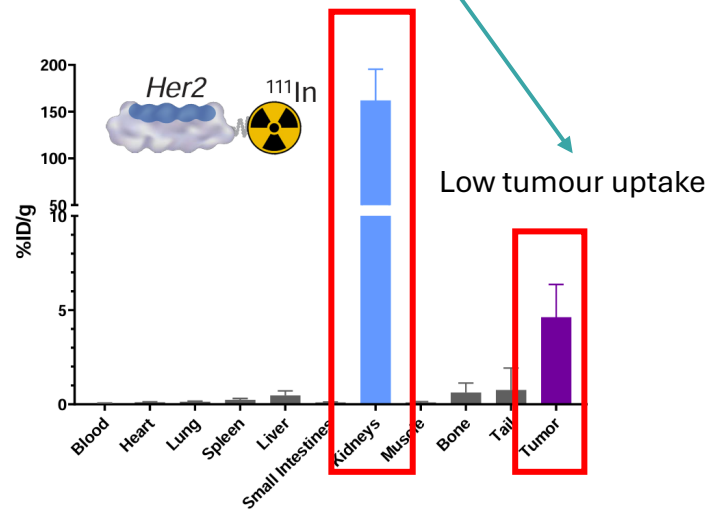


Biodistribution Challenges with Current Approaches:

Tumour:Kidney ratio **highly negative**

High kidney exposure

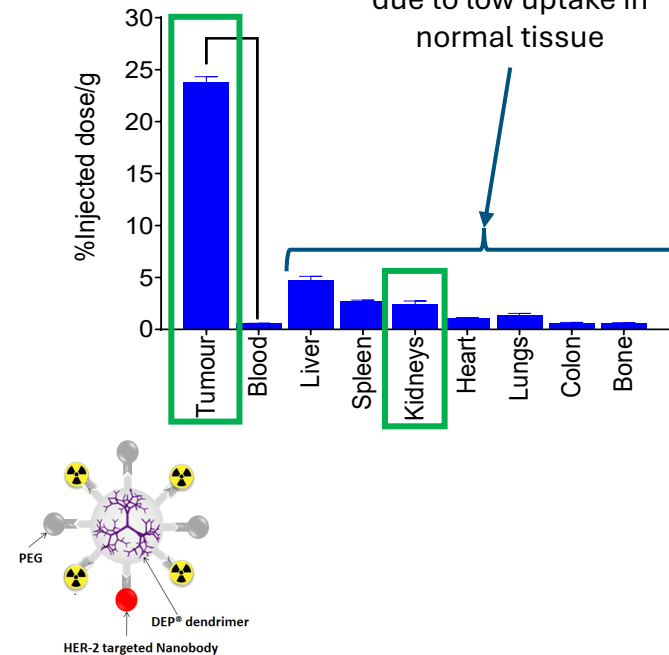
Low tumour uptake



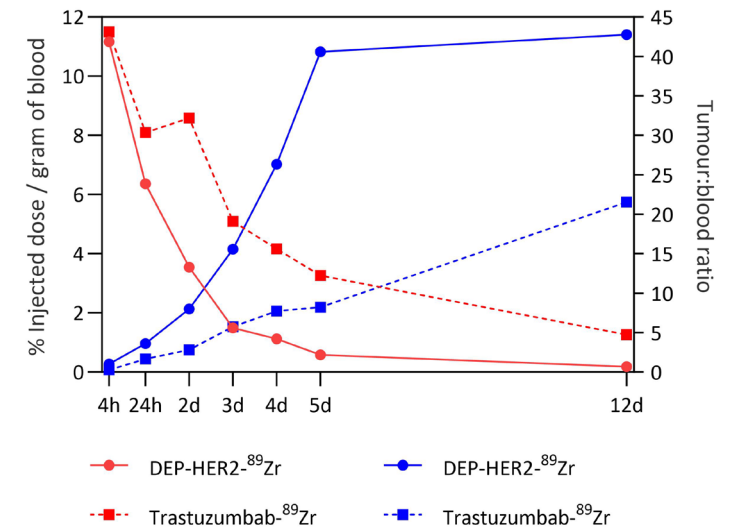
Starpharma's DEP® Technology Solution:

Tumour:Blood ratio >40

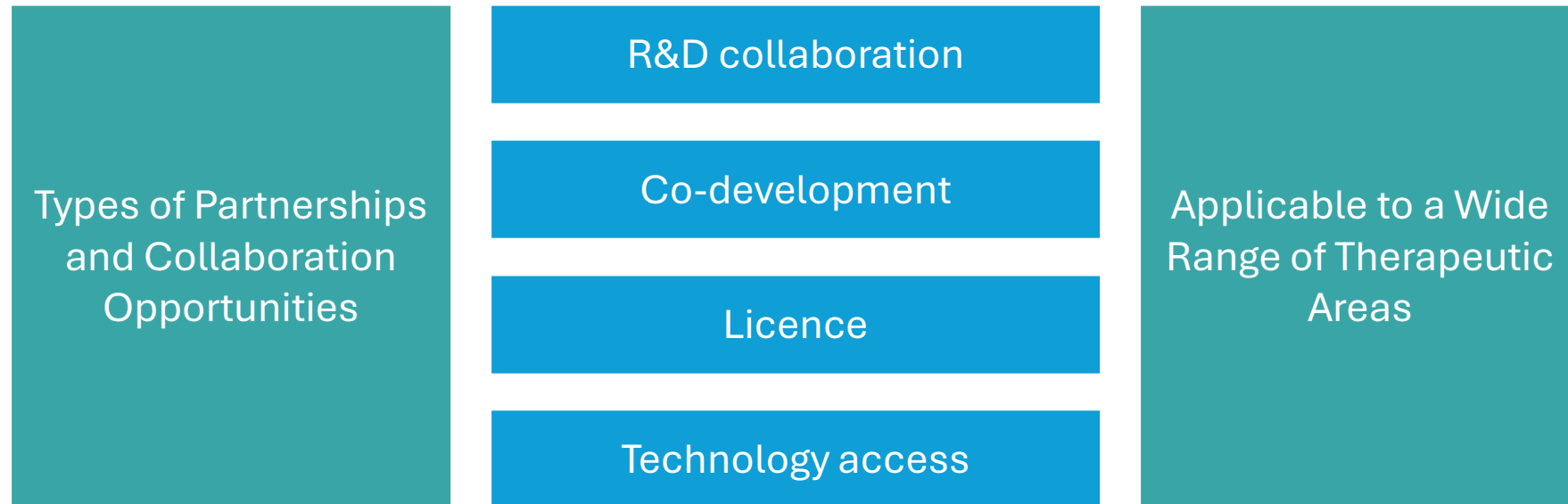
Tumour:Kidney/Organ ratios **highly positive** due to low uptake in normal tissue



DEP® HER2 radiodiagnostic: faster clearance from blood and sustained elevated tumour: blood ratios compared to trastuzumab-⁸⁹Zr



Extensive Partnership Experience, Broad DEP[®] Application Opportunity and a Flexible Approach to Collaboration



Partnerships

Genentech
A Member of the Roche Group

medicxi

 **MSD**

Well Positioned to Accelerate Product Pipeline and Strategic Growth Initiatives

3-Year Summary	FY24 \$M	FY23 \$M	FY22 \$M
Revenue and other income	9.8	4.3	5.2
Loss for the period	(8.2)	(15.6)	(16.2)
Net operating cash outflows	(7.0)	(14.3)	(13.2)

Multiple Revenue Streams

- Licenses and Milestone Payments
- Marketed Products: VivaGel[®] BV and Viraleze[™]
- R&D Income

Cash at 30 September 2024: \$24.0M

Recent Strategic Review Confirmed Three Key Focus Areas to Optimise Shareholder Returns

01

Maximise DEP[®]
asset value

*Prioritising DEP[®] SN38 and DEP[®]
cabazitaxel*

02

Accelerate early
asset
development

*Advancing DEP[®]
radiopharmaceuticals and
partnerships*

03

Build long-term
sustainability

*Increasing revenue,
strengthening IP position, and
fostering a high-performance
culture*

Catalysts to Anticipate in The Next 12 Months

Poised for Value Creation

Over 20 years of experience in advancing dendrimer technology from the lab to the patient.

Clinical and commercial validation of platform technology.



1) License / collaboration for a DEP[®] asset to commercialise



2) Radiodiagnostic progress to the clinic



3) Strategic partnership – expansion and/or licence



4) New VivaGel[®] BV EU Partner



5) Increasing revenue contributing to sustainability



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Thank you.

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