



R&D Marketing Interface

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Learning Objectives

- Highlight the importance of commercial team input into product development strategy in a early stage biotech/medical device company
- Understand the role of marketing in early stage companies to shape the product life cycle to achieve the best commercial success.
- Describe the process of creating a commercially appealing target product profile (TPP).
- Learn how to develop a TPP framework to deliver better outcomes.





Trends

Health care priority: cost control, access and price transparency

Consumerism will influence healthcare decisions

Decreasing access to physicians

Shifting power balance toward payers

Reimbursement models focus on value/outcome

Growth of alternative delivery models and partnerships

Digital transformation enhancing patient centricity and engagement



DEVELOPING A NEW MEDICINE TAKES AN AVERAGE OF 10-15 YEARS



Sources: Drug Discovery and Development: Understanding the R&D Process, www.innovation.org; CBO, Research and Development in the Pharmaceutical Industry, 2006, Tufts Center for the Study of Drug Development, Impact Report May/June 2018, Clinical Development Success Rate 2006-2015, Biotechnology Industry Organization, 2006, Wong C.H., Siah K. W. "Estimation of Clinical Trials Success Rates and Related Parameters, Biostatistics, 20: 273-286, 2019





New Active Substance (NAS) Launched in US 2008-2017



Source: Medicines use and spending in the U.S.- IMS Report, April 2018





Pharma Loss of Exclusivity

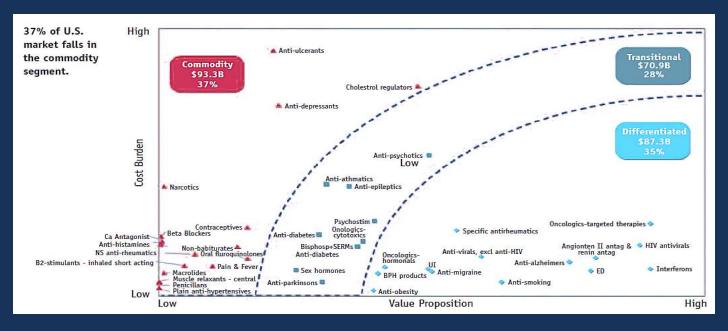








Segments of Pharmaceutical Market



Source: Understanding the New Commercial Models in the Pharmaceutical Industry – An IMS Report, 2009





Brands vs. Generics Shares

2012	2013	2014	2015	2016
317.8	331.5	378.5	425.3	450.0
71.6%	71.2%	71.8%	72.9%	74.2%
16.3%	16.7%	17.1%	16.5%	15.0%
12.2%	12.1%	11.0%	10.6%	10.8%
	317.8 71.6% 16.3%	317.8 331.5 71.6% 71.2% 16.3% 16.7%	317.8 331.5 378.5 71.6% 71.2% 71.8% 16.3% 16.7% 17.1%	317.8 331.5 378.5 425.3 71.6% 71.2% 71.8% 72.9% 16.3% 16.7% 17.1% 16.5%

Branded 15.9% 13.5% 12.2% 11.3% 10.5% Unbranded Generic 77.8% 80.5% 82.1% 83.4% 84.6% Branded Generic 6.3% 5.9% 5.6% 5.3% 4.9%

Source: Medicines use and spending in the U.S.– IMS Report, May 2017



CMS Framework Categorizing Payments to Providers

O2019



Fee for Service – No Link to Value

Payments are

Category 2:

Fee for Service – Link to Quality

Category 3:

Alternative Payment Models Built on Fee-for-Service Architecture

Category 4: Population-Based Payment

Description

- based on
 volume of
 services and not
 linked to quality
 or efficiency
- At least a portion of payments vary based on the quality or efficiency of health care delivery
 - Some payment is linked to the effective management of the population or an episode of care
 Payments still triggered by delivery
 - of services, but opportunities for shared savings or 2-sided risk
- Payment is not directly triggered by service delivery so volume is not linked to payment
- Paid and responsible for the care of a beneficiary for a long period (e.g., ≥1 year)

Medicare Fee-for-Service examples for-service

Majority of
Medicare
payments now
are linked to
quality

Medicare fee-

Limited in

- Hospital valuebased purchasing
- Physician Value Modifier
- Readmissions / Hospital Acquired Condition
 Reduction Program

- Accountable Care Organizations
- Medical homes
- Bundled payments
- Comprehensive Primary Care initiative
- Comprehensive ESRD
- Medicare-Medicaid Financial Alignment Initiative Fee-For-Service Model

- Eligible Pioneer Accountable Care Organizations in years 3-5
- Maryland hospitals

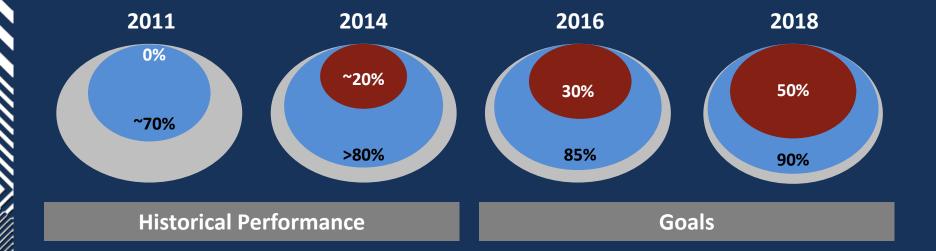




CMS Target Percentage of 'Alternative Payment Models' by 2016 - 2018



Alternative payment models (Categories 3-4) FFS linked to quality (Categories 2-4) All Medicare FFS (Categories 1-4)







Innovative Product

A differentiated product (solution) that offers a meaningful advantage (value) over existing treatments for a given condition

How can marketing shape the product?





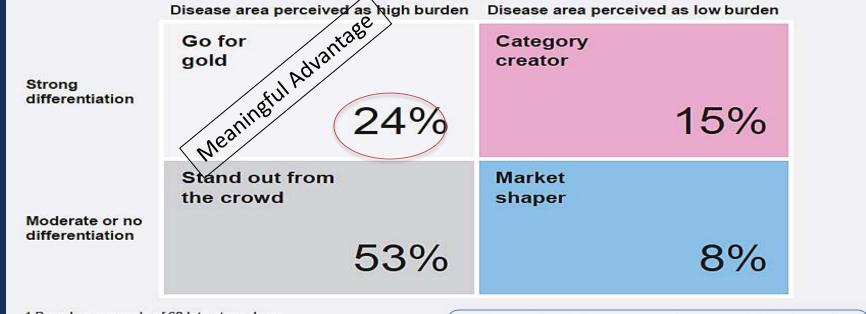
Objectives, 4 P's, A's of Marketing

Objectives	4Ps	4As
Address Unmet Needs	Product	Acceptability
Value to Payers	Price	Affordability
Create Convenience	Place	Accessibility
Communication of Value	Promotion	Awareness





What Proportion of the Launches are Innovative?



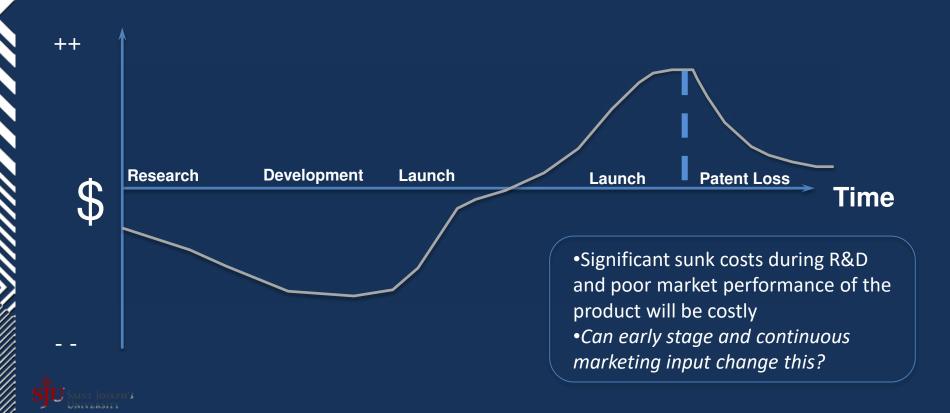
1 Based on a sample of 60 late-stage drugs Source: Evaluate; McKinsey analysis

In the value driven post Accountable Care Act (ACA) era only 24 percent of the launches would be truly innovative



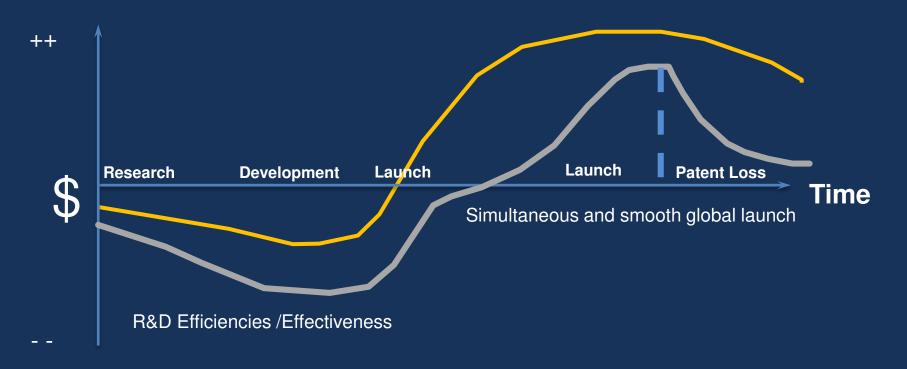


How to Improve the Probability of Success?





Ideal Product Life Cycle







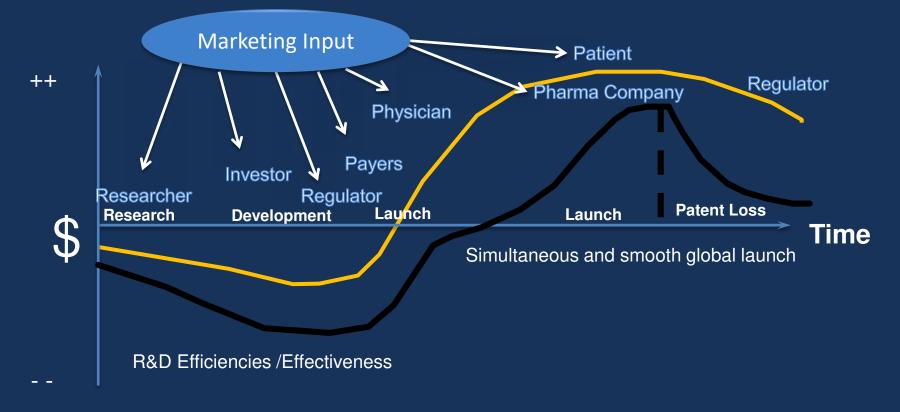
Meeting Stakeholders Needs





Early Marketing Input Can Improve Product Success

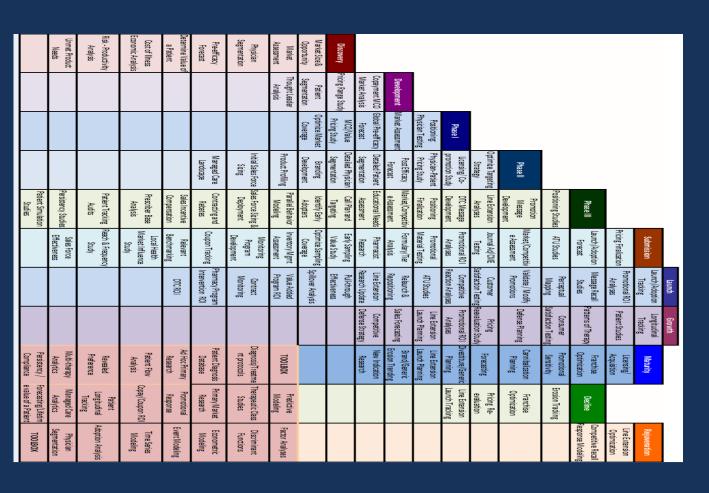






Early Marketing Input Can Improve Product Success









What will Collaboration Achieve...? From....

Stages of Sevelopmen

Discovery

Exploratory Development

Full Development

Registration

Launch

Post-Launch Growth

Organizational Roles

Discovery

Clinical

Regulatory

Manufacturing

Sales

Pre-Clinical

Marketing

IP

HEOR/Pricing/Reimbursement





What will Collaboration Achieve...? To....an Integrated Model

Organizational Roles

Discovery

Exploratory Development Full

Development

Registration

Launch

Post-Launch Growth



Upstream Role of Marketing is Critical for developing an Ideal Label and Product Success!





Marketing Facilitate Cross-Functional Decisions





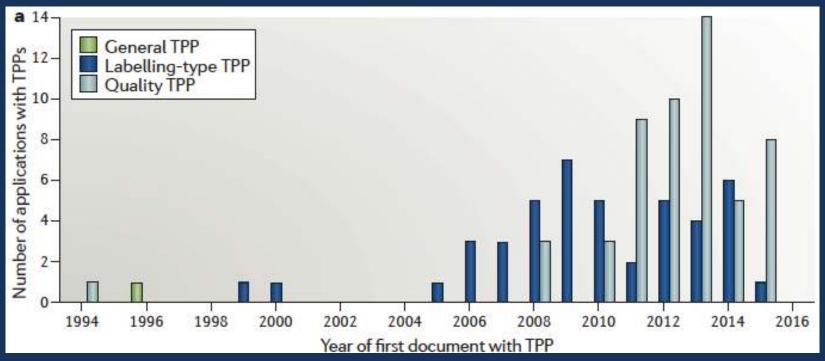
Source: Jambulingam, T. (2018), The R&D Marketing Interface in BioPharma and MedTech, Journal of Commercial Biotechnology, 24(1), 48-55.



What is TPP?

- In 2007 FDA created a guidance document on TPP as a strategic process development tool to facilitate effective communication between the industry and review staff
- TPP is the directional tool that has a significant impact on the drug development process and in particular, its marketing organization
- TPP convert discoveries into companies!
- Marketing as part of commercial team can shape the TPP

Research shows TPP is valuable but underused

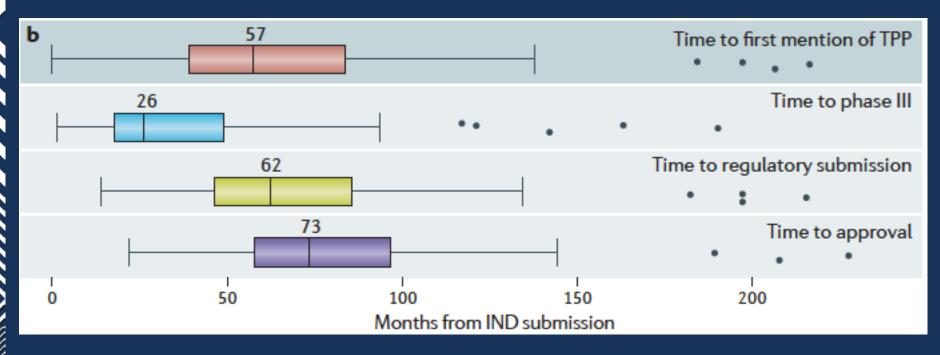


Source: Tyndall et.al. The TPP as a tool for regulatory communication: advantageous but underused, Nature, March 2017, pp. 156





Research shows TPP Valuable but Underused

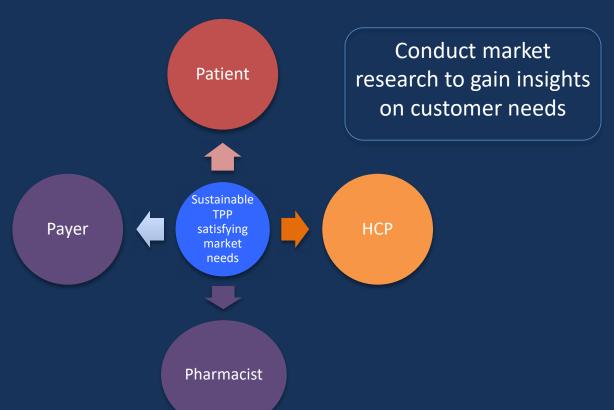


Source: Tyndall et.al. The TPP as a tool for regulatory communication: advantageous but underused, Nature, March 2017, pp. 156





Start with end in mind: How should the label look to meet customer needs?





How can TPP be shaped by Marketing?



<u>TPP</u>

- Indication
- Dosage form
- Dose, frequency
- Differentiation
 - Efficacy
 - Safety
 - Economic



Attributes Shaped by Marketing

- Proposed indication
- Develop formulation
- Develop trade-dress
- Establish efficacy/superiority
- Establish safety advantage
- Develop for pediatric use
- Pharmacoeconomic data





superior clinical

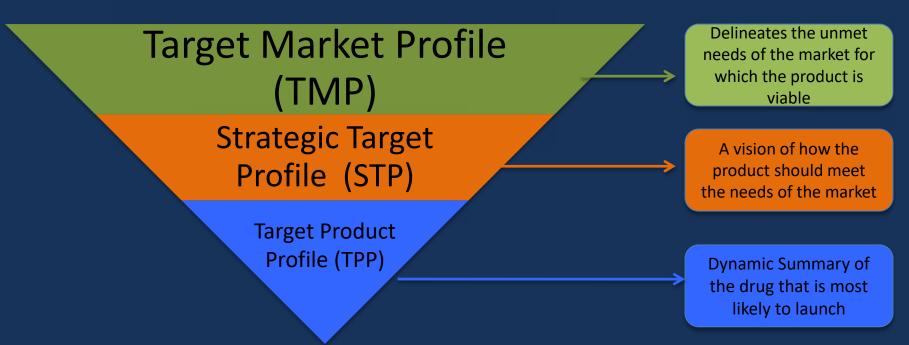
economic value

Who should work together?





STRATEGIC FRAMEWORK



Source: Tebbey, Paul W. and Charles Rink, "TPP: A Renaissance for its Definition and Use, Journal of Medical Marketing, Vol. 9 (4), 301-307.



Strategic Framework



	Target Market Profile (TMP)	
Purpose	Captures all the key information about the market	
Content	 Therapeutic areas/diseases Unmet Need Patient Populations Drivers of use Competitive assessment Economic cost of disease 	
Rigidity	Create before the STP or TPP Details are updated as findings emerge, but core facts change only in response to major market events	

Strategic Framework



	Target Market Profile (TMP)	Strategic Target Profile (STP)	Target Product Profile (TPP)
Purpose	Captures all the key information about the market	A vision for a product that will meet the needs of the market	✓ Positioning
Content	Therapeutic areas/diseases • Unmet Need • Patient Populations • Drivers of use • Competitive assessment • Economic cost of disease	Target attributes (desired profile) Value drivers/Positioning Global Reach Pricing/Reimbursement Patient Share Revenue – Profitability Pharmacoeconomics Investments (R&D, COGS, SGA) Cost of goods Licenses, Royalties	✓ Global Sales Forecast ✓ Developmental Logic ✓ Regulatory and Reimbursement Strategy ✓ Product Value
Rigidity	Create before the STP or TPP Details are updated as findings emerge, but core facts change only in response to major market events	Set at the beginning of clinical development and updated only when necessitated by changes in the TMP	



Strategic Framework

	Target Market Profile (TMP)	Strategic Target Profile (STP)	Target Product Profile (TPP)
Purpose	Captures all the key information about the market	A vision for a product that will meet the needs of the market	A record of the drug that is most likely to launch
Content	Therapeutic areas/diseases • Unmet Need • Patient Populations • Drivers of use • Competitive assessment • Economic cost of disease	Target attributes (desired profile) • Value drivers • Global • Pricing/Reimbursement • Patient Share • Revenue – Profitability • Pharmacoeconomics • Investments (R&D, COGS, SGA) • Cost of goods • Licenses, Royalties	Indications and usage (label) Dosing and administration Contraindications Warnings and precautions Adverse reactions Description Clinical Pharmacology Clinical Studies Storage and handling
Rigidity	Create before the STP or TPP Details are updated as findings emerge, but core facts change only in response to major market events	Set at the beginning of clinical development and updated only when necessitated by changes in the TMP	Updated as clinical and pharmacologic findings emerge and in response to guidance from regulatory authorities

Big

Questions that needs to be asked and and answered during the TPP process

- What is the product description?
- What data or literature is available for review for the various indications and claims?
- What is the unmet need, clinical benefit or value to others?
- Will the product be used for a new or existing procedure?
- What is the standard of care (SOC) for this indication?
- What is the future direction of SOC?
- What is the market potential for each indication and claim?
- What is the probability of success for each indication and claim?
- What are the product's possible differentiating features and will they be obsolete in 5 years?
- What are all of the possible indications for this product (neurovascular, pulmonary, peripheral vascular, gastrointestinal, etc.)?

- What are all of the possible differentiating claims?
- Can premium pricing be justified?
- If so, will payors directly reimburse?
- How is the competitor successful?
- Where does the competition fall short?
- Does IP exist or can it be created?
- Can exclusivity be achieved with a more complex regulatory or clinical strategy?
- If so, what is the company's tolerance or resource availability for such complexity?
- What are the COGS?
- How do development costs compare against five-year return on investment (POI)?
- How does the net present value (NPV) or ROI compare against other projects?

Source: Begin with End in Mind – White Paper Premier Research, 2015



Portfolio Optimization – Go/No Go

Specification – TPP – Current, Minimal, Ideal & Expected

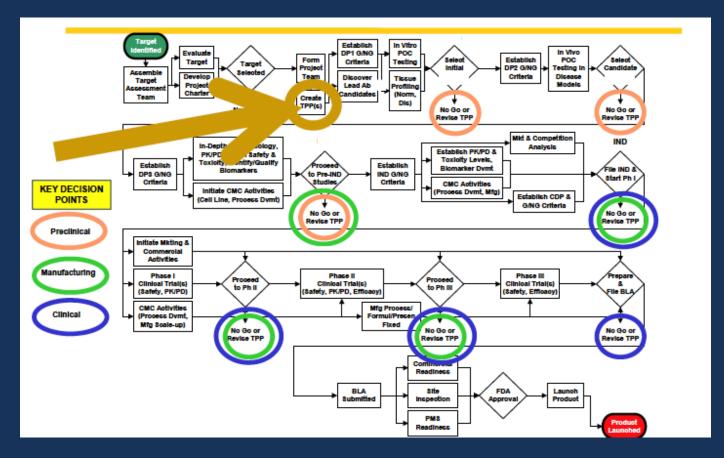
Resources – Manpower and Cost

• Timeline – Milestone Schedule

Risk – Probability of Success (Technical, Commercial)



TPP in Go/No Go Decisions







TPP Criteria

	Current	Minimum	Ideal	Expected
Efficacy One	52%	33%	75%	60%
Efficacy Two	80%	75%	95%	84%
Safety One	67%	60%	85%	82%
Safety Two	45%	50%	30%	35%
Convenience	25%	20%	40%	30%
Cost	\$XXXXX	\$YYYYY	\$ZZZZZ	\$AAAAA





Sample TPP

Product Properties	Minimum Acceptable Result	Ideal Result
Primary Indication	Relief of pain symptoms in diabetic neuropathy	Relief of symptoms in neuropathic pain syndromes
Patient Population	Adults with diabetes who experience neuropathic pain	Adults and children with neuropathic pain
Treatment Duration	Chronic	Chronic
Delivery Mode	Oral	Oral
Dosage Form	Tablet or capsule	Tablet or capsule
Regimen	1-2x/day	1x/day
Efficacy	A 40% decrease in pain score in 30% of patients	A 70% decrease in pain score in 50% of patients.
Risks/Side Effects	Devoid of opioid side effects Devoid of GI side effects from Non-steroidal anti-inflammatory drugs (NSAIDs) Minor or moderate CNS side effects	Devoid of opioid side effects Devoid of GI side effect from NSAIDs No CNS side effects

Source: https:///neuroscienceblueprint.nih.gov/resources/target-product-profile.htm. accessed June 14, 2017



Best Development Strategies...

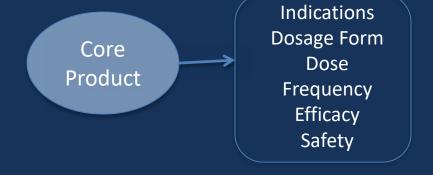
- Use the strategic framework (TMP, STP) to shape TPP and define clinical and commercial value
 - TPP provides developmental logic and saves cost to drug discovery and development program and meet the needs of the market place
 - Encourages right dialog within the company and with the FDA to optimize label and promotability for commercial success

The ideal development strategy identify key milestones -critical times, when the ability of a project to attain its TPP can be assessed - and establish "go / no go" success criteria



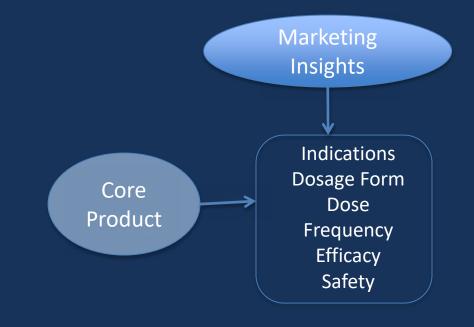








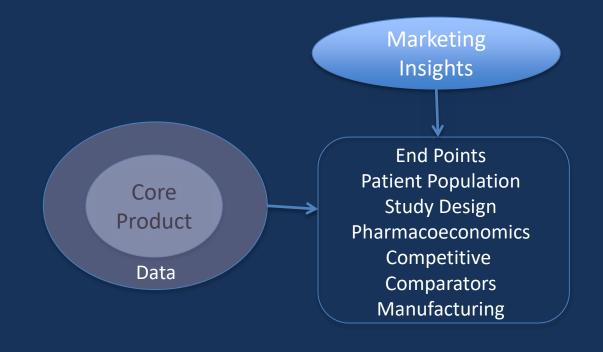




Marketing should shape the "Label" for the product

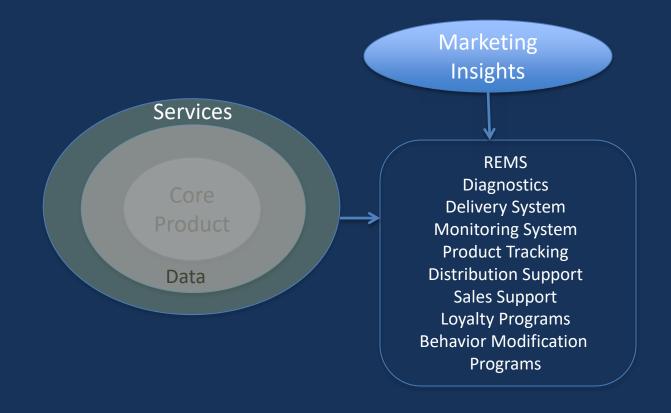






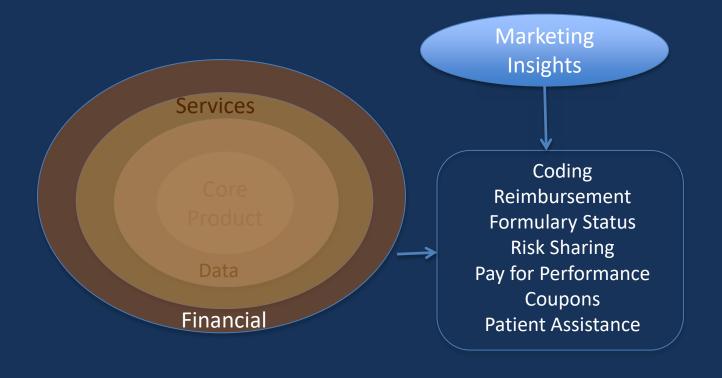






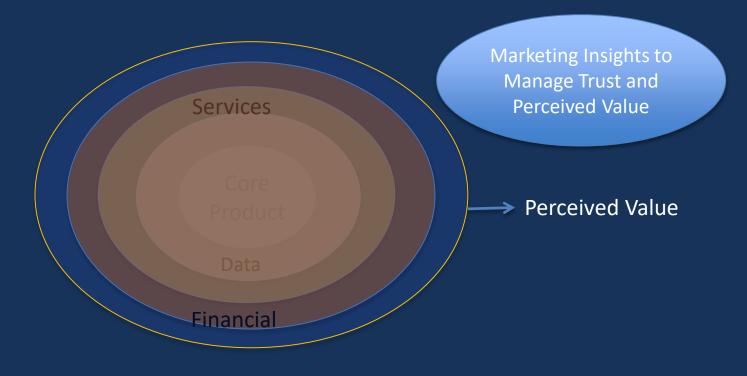
















Final Remarks

- Start with end in mind
- Strengthen the R&D Marketing (Commercial) interface
- Assemble cross functional commercial development team
- Assign a marketing manager to the development team
- Incorporate market research and competitive intelligence in clinical trial planning and label development
- Engage payers early on to get valuable input in development
- Success is when the final version of TPP is similar to the annotated draft labeling!





Questions?

