

2005 Merrill Lynch Japan Conference "A Global Power"

### 'Value Creation; Eisai Way'

### Eisai Co., Ltd.

September 7, 2005

## **Patient Value**

#### **Fulfillment of unmet medical needs**

### Stable supply of quality products

# Provision of pharmaceutical product safety and efficacy information

## **Pipeline Policy**

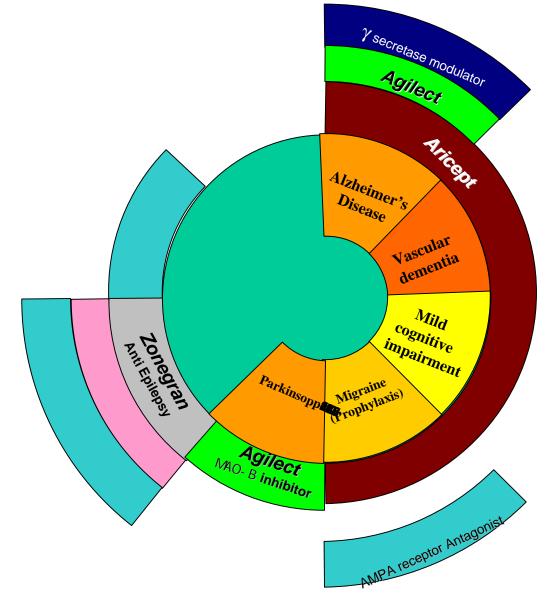
- 1. Area Focus Neurology GI Oncology and Critical Care
- 2. First-in-class Drug
- 3. Active Business Development

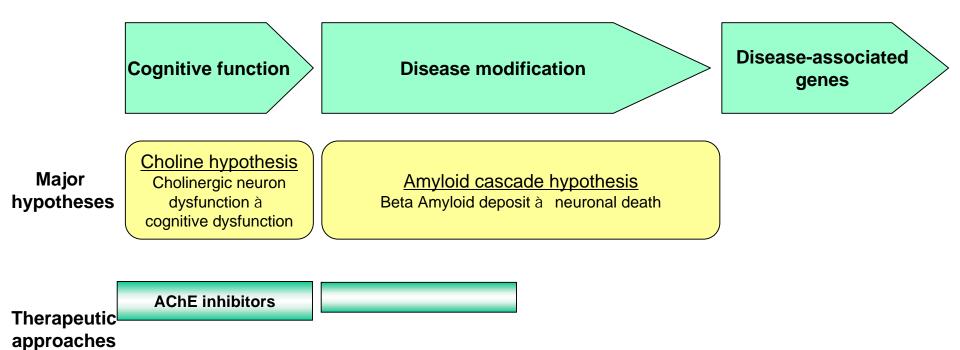
#### Stage

#### Filed for approval

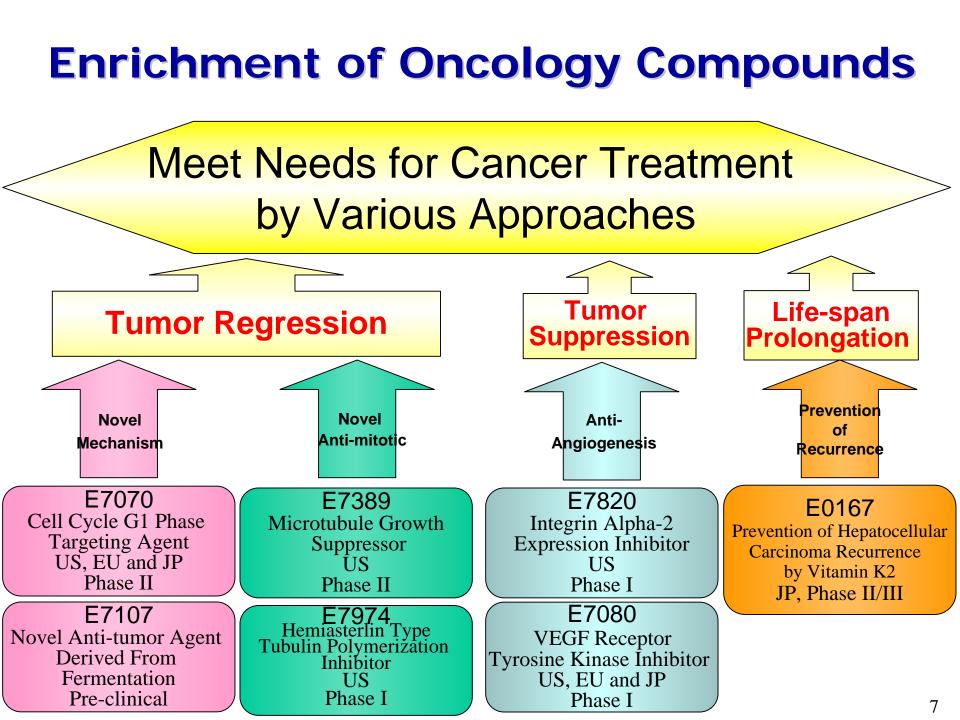
Phase III	Sibutramine (Obesity Management)	Serotonin/Noradrenarin reuptake inhibitor	Natural body weight loss based on dual actions of appetite suppression and energy consumption increase
Preparing Phase III	E2007 (Parkinson's disease)	AMPA receptor antagonist	Reduction of off-time in PD as adjunct therapy with levodopa Excellent safety profile; no worsening of dyskinesia
	E5564 (Sepsis)	Endotoxin antagonist	Reduce mortality and morbidity; good safety profile
	E7389 (Cancer)	Microtubule growth suppressor	Better anticancer efficacy than taxanes Good tolerability, less neurotoxicity
Phase II / III	Adalimumab (Rheumatoid arthritis)	Anti-TNF antibody	Strong and long-lasting efficacy for RA symptoms
	E0167 (Hepatocellular carcinoma)	Vitamin K <sub>2</sub>	Reduce recurrence of hepatocellular carcinoma
	E2014 (Cervical dystonia)	Botulinus toxin	Effective in patients resistant to existing Botulinus toxin
Phase II	E7070 (Cancer)	G1 Phase targeting	Different anticancer spectrum from existing cytotoxics
Phase I	E5555 (Prevention of major cardiac events)	Thrombin receptor antagonist	Dual action of anti-platelet and smooth muscle cell proliferation inhibition
	E7820 (Cancer)	lpha2 integrin suppressor	Survival benefit due to chronic tumor growth suppression
	E7080 (Cancer)	VEGFR kinase inhibitor	Survival benefit due to chronic tumor growth suppression
	E7974 (Cancer)	Tubulin inhibitor	Effective in multi-drug-resistant tumors

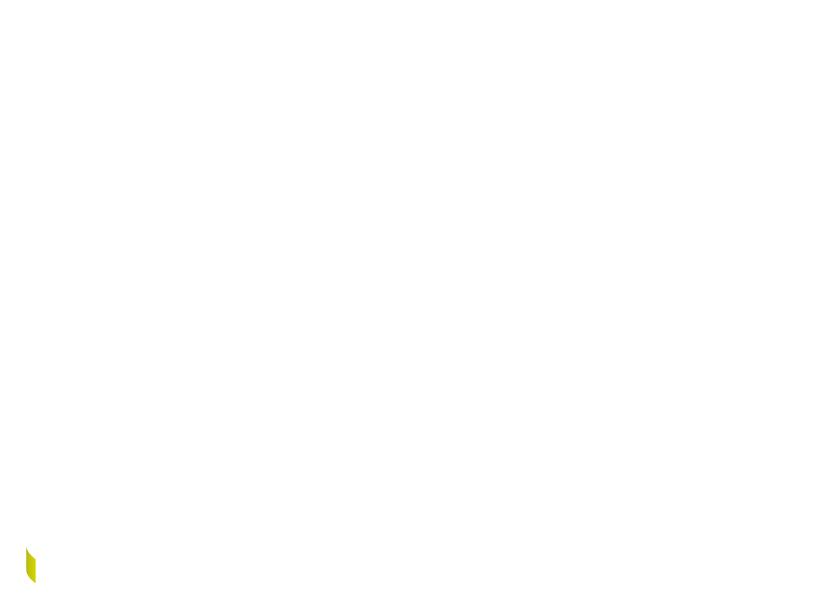
### Fulfilling Pipeline in Neurology Area with Focus on Neurodegenerative Disease

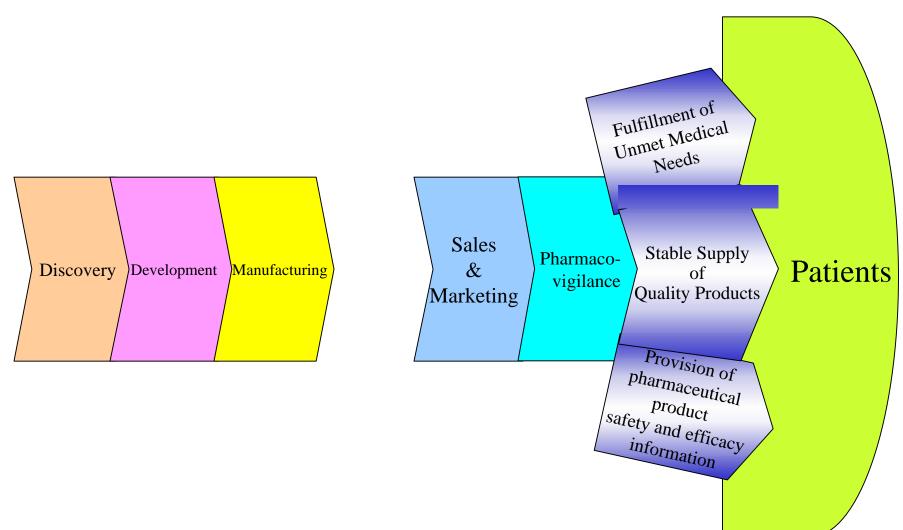












### **Reinforcing Seamless Value Chain**

- ¥ Reconfiguration of P-1 plant in Kashima for preparing drug substance of E7389
- ¥ Construction of the second production facility in Suzhou Plant (China) corresponding to increasing sales in China
- ¥ Expansion of Eisai Research Institute of Boston, Inc. (US)



Eisai Research Institute of Boston, Inc.

## **Shareholder Value**



#### **Return to shareholders**

#### **Investor Relations**

### Growth

#### 1. Pipeline

- E2007 : AMPA Receptor Antagonist
- E7389 : Microtubule Growth Suppressor
- E5564 : Endotoxin Antagonist
- E5555 : PAR-1 Receptor Antagonist
- E2012 : γ Secretase Modulator

POC Success POC Success POC Success Phase I Pre-Clinical

#### 'All First-in-class Type of Drugs'

- 2. Global Opportunities (FY2000 FY2004)
  - Overseas Net SalesCAGR 21.7%Overseas Operating Income\*CAGR 37.9%Consolidated Net SalesCAGR 10.2%Consolidated Operating IncomeCAGR 10.2%

**'49.7% of Net Sales, 59.0% of Operating Income Attributed to Overseas'** 

3. Strong Current Products (FY2000 - FY2004)
Aricept CAGR 23.0%
Aciphex/Pariet CAGR 24.7%

#### 'Active New Indications/Formulations Research'

\* Operating income before Royalty deduction

### Proof of Concept Success -Three First-in-class compounds-

#### • E2007: Oral AMPA receptor antagonist

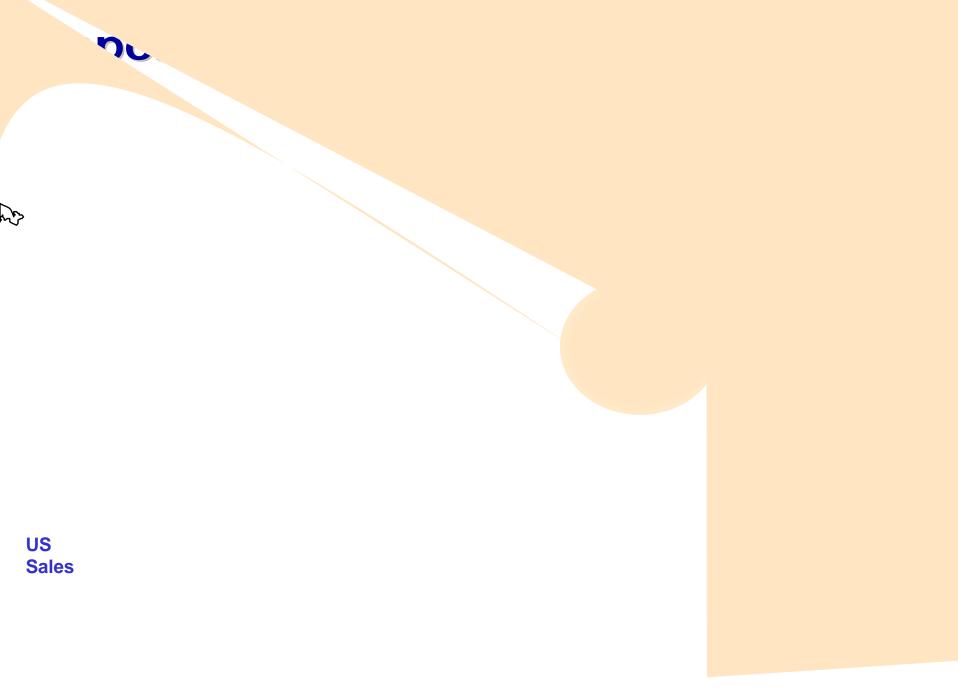
- A placebo-controlled Ph IIb study in patients with Parkinson's disease demonstrated clinically meaningful efficacy (OFF time reduction) and good safety
- Target NDA/MAA in 2Q FY2007
- Ph IIb studies ongoing in migraine prophylaxis and epilepsy, in preparation for multiple sclerosis

#### E7389: Microtubule growth suppressor

- Considerable objective tumor responses were observed in Breast and Non-small cell lung cancers. – those patients had been treated with chemotherapies including taxanes
- No severe neurotoxicities have been observed
- Aiming Subpart-H submission in FY2006

#### • E5564 (eritoran): Endotoxin antagonist

- A placebo-controlled Ph IIb study in patients with severe sepsis showed more than 5% decrease in mortality in high-dose E5564-treated group, with clear dose response
- Good tolerability was demonstrated
- Global Ph III study to be initiated in FY2005
- Target NDA/MAA in FY2008



# **'Value Creation Leads to Enhancement of Eisai's Enterprise Value'**