Information Meeting

March 7, 2014 Eisai Co., Ltd.

Safe Harbor Statement



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- Forecast or target figures in this material are not official earnings guidance but present the midterm strategies, goals, and visions. Official earnings guidance should be referred to in the disclosure of the annual financial report (*kessan tanshin*) in accordance with the rules set by Tokyo Stock Exchange.
- Materials and information provided during this presentation may contain so-called "forward-looking statements." These statements are based on current expectations, forecasts and assumptions that are subject to risks and uncertainties which could cause actual outcomes and results to differ materially from these statements.

Modern Pharmaceutical Industry Environment Factors



- Disease Commonness: enlarging
- Regulatory Approval: efficient
- Payors: powerful
- Pricing & Reimbursement: complication
- International Reference Pricing: expansion
- Medical Practice Standardization: progression

Expansion of global business opportunity

FDA, EMA, and PMDA



Transition of number of approvals and length for review

		Number of approvals*1					Average length for review*2 (median) (months)				
		2008	2009	2010	2011	2012	2008	2009	2010	2011	2012

FDA

Eisai's Policy Framework





1. hhc

2. Innovation

3. Access

Shift from Two Brand Company to Multi-Brand Company



Two big brand company (Aricept and Pariet/AcipHex)



Multi-Brand Company



Simultaneous establishment of each brand in all marketplaces is the key for success



Necessity of changing business model

Background for Business Unit Transformation



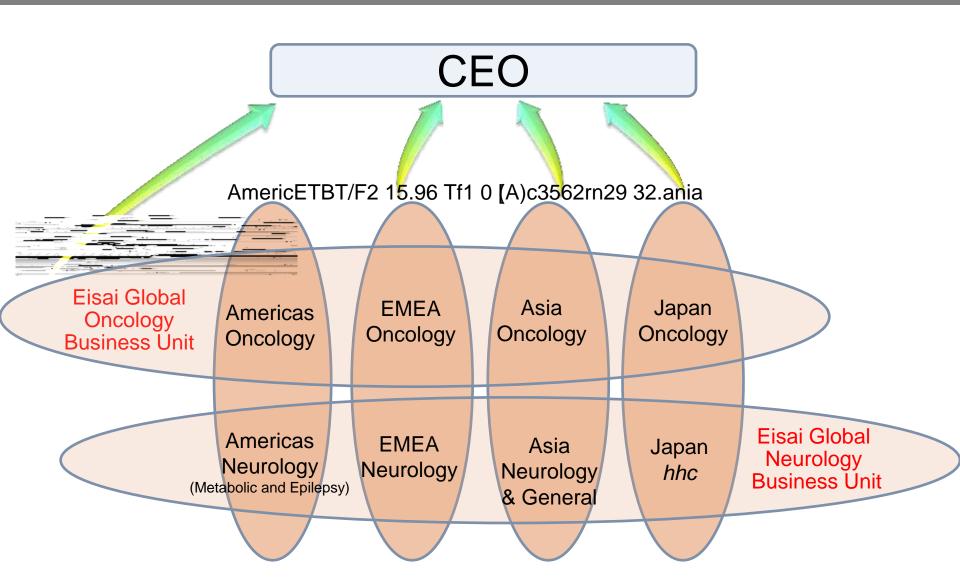
- Generalization of simultaneous global development
 - Connection of global markets in pricing/ reimbursement
 - Standardization of medical practice

Initiation of international collaboration between agencies and reimbursement agencies

Risk of volatile operation under different strategies in each region

Necessity to operate under integrated global brand strategy based on each region s circumstance





Purpose for Establishment of Global Business Unit



 Global Business Unit creates integrated brand strategy based on knowledge and experience in each region

Business units in each region are essential and important parts of the Global Business Unit

Global Brands with Great Potential





- Late line treatment for metastatic breast cancer
- Earlier line treatment for metastatic breast cancer*1 (Europe)
- Non-small cell lung cancer
- Soft tissue sarcoma

Lenvatinib

- Differentiated thyroid cancer
- Hepatocellular carcinoma
- Aim to submit in FY2014 for differentiated thyroid cancer indication

Fycompa

- Adjunctive therapy for partial-onset seizure
- Primary generalized tonic-clonic seizure
- Expansion of launch countries

BELVIQ®

- Obesity
- Smoking cessation
- Expansion of launch countries

Avatrombopag

 Aim to submit in FY2015 for treatment of thrombocytopenia with chronic liver disease patients who will undergo elective surgical or diagnostic procedures

E2006

Aim to submit in FY2016 as an insomnia treatment

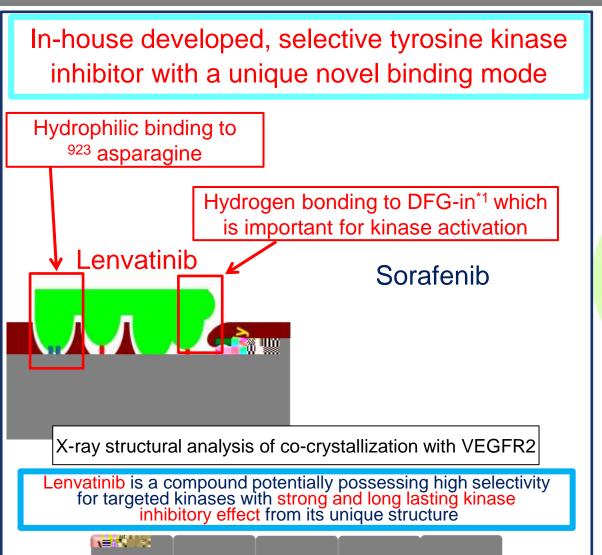
^{*} Indications shown in green are referred for approved indications (lenvatinib is in preparation for submission) and the potential indications are shown in blue
*1: Earlier line treatment for metastatic breast cancer has been approved in Japan



A large-scale global clinical study in breast cancer (Study 301)

Growth Strategy of Investigational Lenvatinib

Selective tyrosine kinase inhibitor with a unique binding mode



Plan global submission and launch of a molecular-targeted small molecule agent with an unique binding mode

Thyroid cancer
Favorable topline
results
Plan simultaneous
submission in
the U.S., EU, and
Japan in FY2014

Hepatocellular carcinoma Phase III study on going in the U.S., EU, Japan, China, and Asia

Global development is underway in various types of cancer

Melanoma

NSCLC*2

11 *2: Non-small cell lung cancer

^{*1:} DFG-in: Conformation where Asp-Phe-Gly (DFG); an important site for activation of kinase, is closed Conformation of DFG-out also exists where DFG is open and binding site has hydrophobic pockets



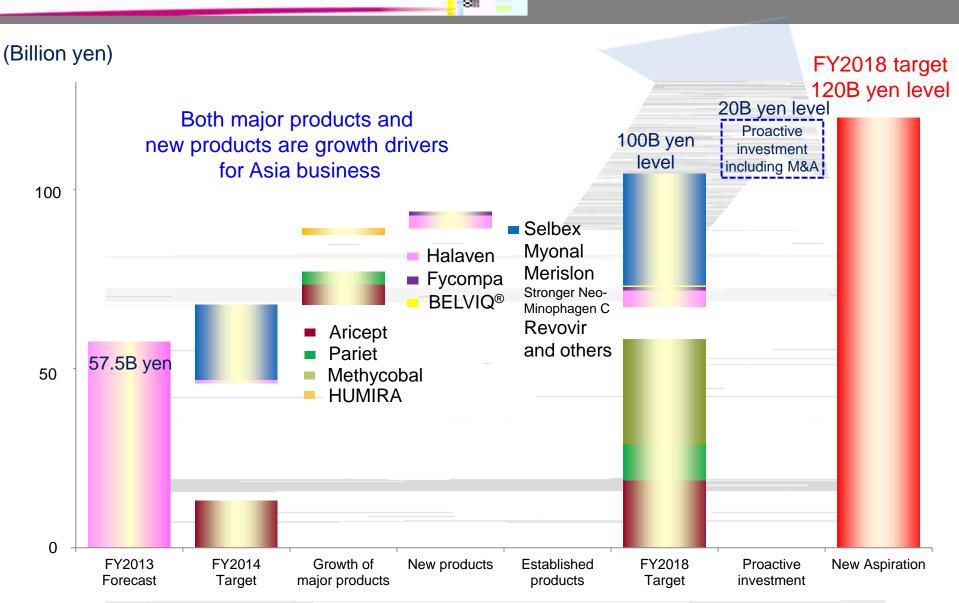
SIII





Trajectory of Asia Business Toward target net sales of 120B yen in FY2018









Launch and Approvals in Strategic Markets Seek patient contribution through independent marketing



	Launch/ Approved	Under review			
Russia	Halaven, Fycompa,	Inovelon			
Mexico		Halaven, Fycompa, BELVIQ [®] , Inovelon, Gliadel, Dacogen, Targretin			
Brazil	Halaven	Fycompa, BELVIQ [®] , Inovelon, Gliadel			
Canada	Halaven, Fycompa,	BELVIQ®			
Australia	Halaven, Zonegran,	Fycompa			
Middle East	Halaven, Aricept, Pariet,	Fycompa			



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• Halaven launched in September 2013

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With Advantages as a Pioneer in the Field of AD Treatment (Various Projects are in Progress



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Strong commitment as a pioneer; developed Aricept,





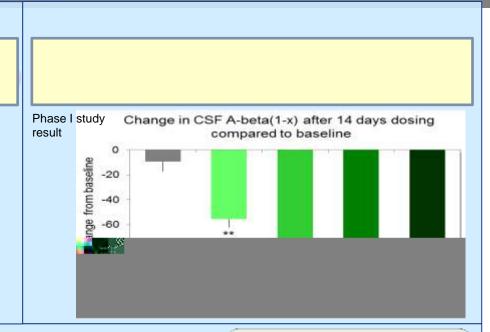
Investigational BAN2401 and E2609



Investigational clinical trial with unique, in-house developed adaptive design



High affinity and selectivity for A-beta protofibrils



POC study

Screening of patients utilizing imaging

Patients with MCI*1 and mild AD with A-beta accumulation

Bayesian adaptive design

196 250 300 350 400 450 500 550

Enables effective assessment of patient arms with higher efficacy and safety

Evaluate with Composite Score developed by Eisai (ADAS-Cog*2, MMSE*2, CDR*2) that enables AD diagnosis in early stage,

clinical progress, and treatment effect

Submission Schedule in Major Countries (NMEs+LCM)



