



# The Experience of a Japanese Company in the UK Market: A View from Eisai

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*hhe*  
human health care

# History of Eisai in the UK



- (Eisai was established in 1941 in Japan)
- 1988 Eisai Laboratories opened on the UCL Campus
- 1988 Eisai UK Ltd was established
- 1995 Eisai UK offices opened in Hammersmith
- 1997 Aricept approved (MCA – UK Reference Member State)
- 1998 Pariet approved



# History of Eisai in the UK, contd



- 2005 EEL (Eisai Europe Ltd) established London HQ
- 2009 EKC opened Hatfield (on schedule, on budget)
- 2010 One Europe Operation introduced
- 2012 Expanded – EMEA + Russia region
- 2013 “Oceania” responsibilities added
- 2013 September – Manufacturing expansion



# Eisai's Business Region Headquarters



EMEA (Europe, Middle East, Africa, Russia & Oceania)

(HQ UK)

HQs

Eisai Japan  
(HQ Japan)

HQs

Asia  
(HQ Japan)

Americas  
(HQ USA)

HQs



- **Lean efficient structure focussed on growing top line sales**



## EMEA Headquarters

- Executive Offices
- Finance & Administration
- Sales & Marketing
- Market Access
- HR, Legal & IP



## Product Creation

- CNS discovery research
- Clinical Development
- Regulatory



## Production

- Solid Dose manufacture
- Packing
- Distribution





# Eisai EMEA Portfolio of Brands



## ONCOLOGY



## EPILEPSY



## HERITAGE BRANDS



# The Selection of the UK as the home for the Eisai European EMEA Knowledge Centre



# Why the UK?



- Science base
- Skilled workforce
- Strong regulatory environment: MHRA and EMA (London is the European centre for medicines regulation)
- Communications, accessibility





# Why the UK? contd

- Strong support from the UK central government, DH, BIS, UKTI, East of England Regional Office, local government
- English language
- Soft factors - driving, international schools, history, feel welcome
- Dialogue with government
- Partnership with government



# Experience since Establishment of EKC



- Continuing excellent dialogue with BIS/DH
- OLS and BIS relationship team – “Buddy Minister”
- MISG dialogue
- NIHR initiatives on clinical research
- MHRA support – scientific advice – (rapporteur)



# Policy Initiatives



- Innovation Report
- Health and Wealth Agenda
- UK as a leader/champion in Europe
- IGFAM – global emerging markets initiative
- NIHR - clinical trials – Scientific Research Agency
- COGENT programme
- MISG manufacturing task force
- Patent Box
- R & D tax credits
- Red Tape Challenge



# Ongoing Issues



- Innovation uptake
- EU regulatory burden
- Reward for innovation, pricing and reimbursement
- HTA (Health Technology Assessment)
- NICE



# The Future

- Global competition for investment
- Emerging markets
- EU Regulatory System – “Lifting the burden” – comparison with other major regulators (FDA/PMDA)
- Evolution of the HTA, need to value and reward innovation
- UK membership of EU
- US/JP/EU trade negotiations



# Concluding remarks

