

Thursday, 3rd October 2013

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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

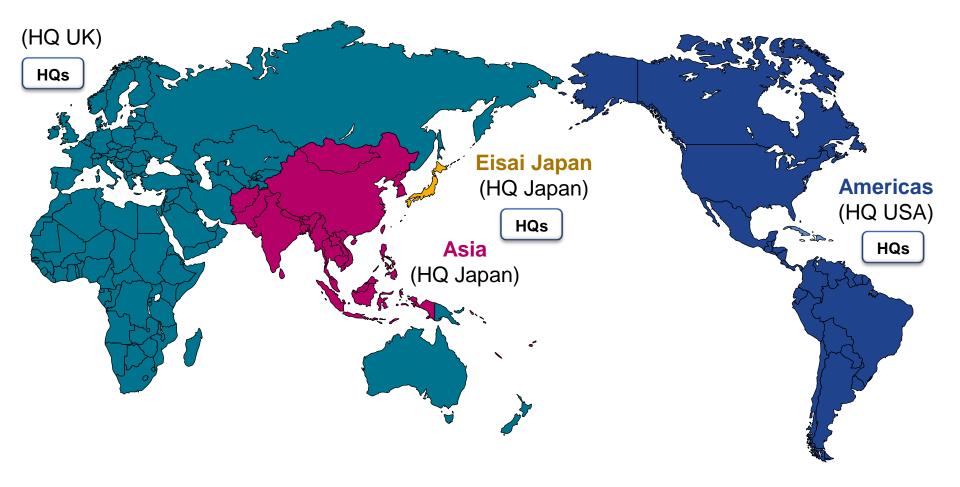




Eisai's Business Region Headquarters



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









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











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



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







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





Eisai

- 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

