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Agendia Postpones its IPO Due to Unfavourable Market Conditions

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Agendia (or the "Company"), a commercial-stage molecular cancer diagnostics company, announces the postponement of its Initial Public Offering ("IPO") that was due to price today. Agendia announced its intention to float on 20 May 2011 and started its book building process on 6 June 2011. The timetable has coincided with an extraordinarily volatile period in global capital markets resulting in high levels of uncertainty and volatility that do not meet the existing shareholders' and the Company's desire to achieve a successful IPO with an orderly aftermarket.

Commenting on this announcement, Bernhard Sixt, Agendia's Chief Executive Officer, said: "We are pleased by the response we have received from investors but current volatile and uncertain market conditions do not allow us to launch the transaction and achieve a smooth transition into the public markets. Agendia's product offering is strong and our world-class team will continue to increase the pace of commercial growth as a private company, supported by our existing shareholders who are committed to the successful future growth of the Company. We would like to thank both Dutch and international investors who took the time to meet us and who showed great interest in our company."

About Agendia

Agendia is a commercial-stage molecular diagnostic company focused on the discovery, development and commercialisation of genomic-based diagnostic products to improve the quality of life for cancer patients by providing healthcare professionals with critical information to enable safe and effective personalised treatment. The Company's Symphony(TM) suite of four complementary breast cancer tests, TargetPrint(R), MammaPrint(R), BluePrint(TM) and TheraPrint(R), provides a comprehensive support system for oncologists to determine whether a breast cancer patient is likely to benefit from hormonal therapy, chemotherapy or targeted therapies, saving patients from unnecessary treatments and lowering healthcare costs. Agendia's lead test, MammaPrint(R), currently the only molecular diagnostic breast cancer recurrence test to receive clearance from the US Food and Drug Administration, gives physicians a tool to clearly and decidedly separate "high" risk from "low" risk recurrence in early stage breast cancer patients, thereby better gauging the "high" risk patients' need for chemotherapy. Agendia is advancing a pipeline of new products, which includes a further extension of its breast cancer suite of tests as well as products for colon cancer and lung cancer. The Company's research and development activities are driven by its deep scientific roots and supported by collaborations with leading academic consortia, cancer centres and pharmaceutical companies.

Agendia was founded in 2003 as a spin-off of the Netherlands Cancer Institute and is based in Amsterdam, the Netherlands, and Irvine, California, United States. For more information, please visit http://www.agendia.com.

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In connection with the Offering, one of the Underwriters (the "Stabilising Manager") (or persons acting on behalf of the Stabilising Manager) may over-allot shares or effect transactions with a view to supporting the market price of the shares at a level higher than that which might otherwise prevail. However, there is no assurance that the Stabilising Manager (or persons acting on behalf of the Stabilising Manager) will undertake stabilisation action. Any stabilisation action may begin on or after the date on which adequate public disclosure of the final price of the shares is made and, if begun, may be ended at any time, but it must end no later than 30 days after the date of commencement of trading of the shares.

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Aan de belegging zijn risico's verbonden. De waarde van de aangeboden effecten kan fluctueren. Rendementen uit het verleden zijn geen garantie voor de toekomst. Potentiële beleggers wordt geadviseerd om eerst hun eigen beleggingsadviseur te raadplegen alvorens een beleggingsbesluit te nemen.

For further information, please contact: Agendia Dr. Bernhard Sixt, President & CEO investorrelations@agendia.com International media and investor enquiries M:Communications Mary-Jane Elliott / Emma Thompson Tel: +44(0)20-7920-2345, E-mail: agendia@mcomgroup.com Dutch media enquiries Huijskens Communications Ian Bickerton Tel +31(0)20-685-5955, E-mail: i.bickerton@huijskens.nl

Contact:

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