

15 June 2010

AquaBounty Technologies, Inc.
(“AquaBounty” or the “Company”)

Update on FDA Approval for AquaAdvantage® Salmon

AquaBounty Technologies, Inc. (AIM: ABTX), a biotechnology company focused on enhancing productivity in the aquaculture market, announces progress on their pending New Animal Drug Application (NADA) for AquaAdvantage® Salmon (AAS) from the U.S. Food and Drug Administration.

The Company confirms that it has received two further letters from the Center for Veterinary Medicine (CVM) advising that they have completed their review on sections four and five of the seven sections of the application. These two recent letters indicate acceptance of the AAS data supporting the durability of its genotype and phenotype, showing that the product is stable and unchanged over multiple generations.

The Company believes that the reviews for the remaining two parts of the application are very nearly complete. As reported previously, all technical submissions necessary for the review and approval of the product have been made and acknowledged. Management has worked constructively with CVM reviewers to answer all questions and is confident of a successful outcome in the near future.

Following formal acceptance of the remaining two technical sections, it is expected that CVM will announce the holding of a Veterinary Medical Advisory Committee meeting on AAS as the next step in their formal process for approval of the product.

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