

Expected R&D milestones

Pipeline charts as communicated at
Q4 and Full Year 2019 Results
Meeting dated February 6, 2020

Products	Expected milestones	Timing
cemiplimab ^{(1)(**)}	Pivotal trial read-out in 2L Basal Cell Carcinoma	H1 2020
Sarclisa [®]	U.S. and EU regulatory decisions in 3L Relapsed-Refractory Multiple Myeloma	Q2 2020
Dupixent ^{®(1)(**)}	U.S. regulatory decision in Atopic Dermatitis for 6-11 year-old age group ⁽²⁾	Q2 2020
MenQuadfi [™]	U.S. regulatory decision for ≥ 2-year old age group	Q2 2020
Fluzone [®] QIV HD	EU regulatory decision for ≥ 65-year old age group	Q2 2020
avalglucosidase alfa	Pivotal trial read-out in Late Onset Pompe Disease	Q2 2020
isatuximab	Pivotal trial read-out in 2L Relapsed-Refractory Multiple Myeloma (IKEMA)	Q2 2020
Dupixent ^{®(1)(**)}	Part A readout from pivotal trial in Eosinophilic Esophagitis	Q2 – Q3 2020
sutimlimab	U.S. regulatory decision in Cold Agglutinin Disease	Q3 2020
SAR440340 ^{(1)(**)} (anti-IL33 mAb)	Proof of concept study read-out in Atopic Dermatitis	Q3 2020
SAR439859 (SERD)	Proof of concept study read-out in Breast Cancer (combo, adj.)	H2 2020
Flublok [®]	EU regulatory decision for ≥ 50-year old age group	Q4 2020
MenQuadfi [™]	EU regulatory decision for ≥ 12-month old age group	Q4 2020
Dupixent ^{®(1)(**)}	Pivotal trial read-out in Asthma for 6-11 year old age group	Q4 2020

(1) Developed in collaboration with Regeneron

(2) Granted breakthrough designation and priority review with FDA Decision May 26, 2020

(**) Partnered and/or in collaboration – Sanofi may have limited or shared rights on some of these products

QIV: Quadrivalent Influenza Vaccine; HD: High-Dose