Commission Implementing Decision (EU) 2017/2375 of 15 December 2017 authorising the placing on the market of N-acetyl-D-neuraminic acid as a novel food ingredient under Regulation (EC) No 258/97 of the European Parliament and of the Council (notified under document C(2017) 8431) (Only the English text is authentic)

Article 1

N-acetyl-D-neuraminic acid as specified in Annex I to this Decision may be placed on the Union market as a novel food ingredient for the uses defined and at the maximum levels established in Annex II to this Decision.

Article 2

1 The designation of *N*-acetyl-D-neuraminic acid authorised by this Decision on the labelling of the foodstuffs shall be '*N*-acetyl-D-neuraminic acid'.

2 Food supplements containing *N*-acetyl-D-neuraminic acid shall be labelled in line with the presentation requirements applied under Regulation (EU) No 1169/2011 with a statement that the food supplement should not be given to infants, young children and children under 10 years of age where they consume breast milk or other foods with added *N*-acetyl-D-neuraminic acid within the same twenty four hour period.

Article 3

This Decision is addressed to Glycom A/S, Kogle Allé 4, 2970 Hørsholm, Denmark.

Done at Brussels, 15 December 2017.

For the Commission

Vytenis ANDRIUKAITIS

Member of the Commission

Changes to legislation:
There are currently no known outstanding effects for the Commission Implementing Decision (EU) 2017/2375.