Consultation on the rationalisation of official control shellfish biotoxin and phytoplankton monitoring points in Carlingford Lough

Status: Closed
Date launched: 21 April 2021
Closing date: 2 June 2021

Summary of responses

Northern Ireland

PDF
View Summary of responses: 2021 Carlingford Lough Biotoxin RA Consultation as PDF(Open in a new window) (176.06 KB)

Who will this consultation be of most interest to?

Carlingford Lough shellfish harvesters, enforcement authorities, official control sampling officers, official control testing laboratories and other interested stakeholders.

Consultation subject

A proposal by the Food Standards Agency (FSA) to rationalise the number of official control monitoring points for biotoxin and phytoplankton monitoring in Carlingford Lough and introduce Representative Monitoring Points (RMPs) and Associated Harvesting Areas (AHAs) .

Purpose of the consultation

To seek comments from industry, enforcement authorities, and other interested stakeholders on the proposed rationalisation of biotoxin and phytoplankton monitoring in Carlingford Lough and the implementation of RMPs and AHAs.

Consultation Pack

Northern Ireland

PDF

View Rationalisation of official control shellfish biotoxin and phytoplankton monitoring points in Carlingford Lough as PDF(Open in a new window) (143.12 KB)

How to respond

Responses to this consultation should be emailed or posted to the addresses below:

Email: executive.support@food.gov.uk

Postal address:

Food Standards Agency in Northern Ireland
Executive Support Unit
10a-c Clarendon Road, Belfast
Co Antrim
BT1 3GB

Publication of response summary

Within three months of a consultation ending we aim to publish a summary of responses received and provide a link to it from this page.

You can find information on how we handle data provided in response to consultations in our Consultations privacy notice.

Further information

This consultation has been prepared in accordance with HM Government Consultation Principles. If an Impact Assessment has been produced, this is included in the consultation documents. If no Impact Assessment has been provided, the reason will be given in the consultation document.