## NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

# Single Technology Appraisal

# Ramucirumab for treating advanced gastric cancer or gastro-oesophageal junction adenocarcinoma previously treated with chemotherapy [ID741]

Consultees	Commentators (no right to submit or appeal)
Company         • Eli Lilly (ramucirumab)         Patient/carer groups         • Independent Cancer Patients Voice         Professional groups         • Association of Cancer Physicians         • Cancer Research UK         • Oesophageal Patients Association         • Royal College of Nursing         • Royal College of Physicians         • Royal College of Radiologists         Others         • Department of Health         • NHS England         • NHS Greater Huddersfield CCG         • NHS Wigan Borough CCG         • Welsh Government	<ul> <li>appeal)</li> <li><u>General</u></li> <li>Department of Health, Social Services and Public Safety for Northern Ireland</li> <li>Healthcare Improvement Scotland</li> <li><u>Comparator companies</u></li> <li>Accord Healthcare (docetaxel, fluorouracil, irinotecan, paclitaxel)(Confidentaility agreement not signed, not participating)</li> <li>Actavis UK (docetaxel, irinotecan, paclitaxel) (Confidentaility agreement not signed, not participating)</li> <li>Bristol-Myers Squibb (paclitaxel) (Confidentaility agreement not signed, not participating)</li> <li>Bristol-Myers Squibb (paclitaxel) (Confidentaility agreement not signed, not participating)</li> <li>Hospira UK (docetaxel, fluorouracil, irinotecan, paclitaxel) (Confidentaility agreement not signed, not participating)</li> <li>medac GmbH (docetaxel, fluorouracil, irinotecan, paclitaxel) (Confidentaility agreement not signed, not participating)</li> <li>Mylan (irinotecan) (Confidentaility agreement not signed, not participating)</li> <li>Pfizer (irinotecan) (Confidentaility agreement not signed, not participating)</li> <li>Sandoz (docetaxel, fluorouracil, irinotecan, paclitaxel) (Confidentaility agreement not signed, not participating)</li> <li>Sanofi (docetaxel) (Confidentaility</li> </ul>

## Matrix of consultees and commentators

Matrix for the technology appraisal of ramucirumab for treating advanced gastric cancer or gastro-oesophageal junction adenocarcinoma previously treated with chemotherapy [ID741] Issue date: March 2015

National Institute for Health and Care Excellence

Consultees	Commentators (no right to submit or appeal)
	Teva UK (docetaxel, irinotecan)     (Confidentaility agreement not     signed, not participating)
	<ul> <li><u>Relevant research groups</u></li> <li>Institute of Cancer Research</li> <li>National Cancer Research Institute</li> </ul>
	<ul> <li>Evidence Review Group</li> <li>Kleijnen Systematic Reviews Ltd</li> <li>National Institute for Health Research Health Technology Assessment Programme</li> </ul>
	<ul> <li><u>Associated Guideline Groups</u></li> <li>National Collaborating Centre for Cancer</li> </ul>
	Associated Public Health Groups None

NICE is committed to promoting equality, eliminating unlawful discrimination and fostering good relations between people who share a protected characteristic and those who do not. Please let us know if we have missed any important organisations from the lists in the matrix, and which organisations we should include that have a particular focus on relevant equality issues.

## PTO FOR DEFINITIONS OF CONSULTEES AND COMMENTATORS

## Definitions:

## <u>Consultees</u>

Organisations that accept an invitation to participate in the appraisal; the company that markets the technology; national professional organisations; national patient organisations; the Department of Health and the Welsh Government and relevant NHS organisations in England.

The company that markets the technology is invited to make an evidence submission, respond to consultations, nominate clinical specialists and has the right to appeal against the Final Appraisal Determination (FAD).

All non-company consultees are invited to submit a statement<sup>1</sup>, respond to consultations, nominate clinical specialists or patient experts and have the right to appeal against the Final Appraisal Determination (FAD).

#### **Commentators**

Organisations that engage in the appraisal process but that are not asked to prepare an evidence submission or statement, are able to respond to consultations and they receive the FAD for information only, without right of appeal. These organisations are: companies that market comparator technologies;

Healthcare Improvement Scotland; the relevant National Collaborating Centre (a group commissioned by the Institute to develop clinical guidelines); other related research groups where appropriate (for example, the Medical Research Council [MRC], National Cancer Research Institute); other groups (for example, the NHS Confederation, NHS Alliance and NHS Commercial Medicines Unit, and the *British National Formulary*.

All non-company commentators are invited to nominate clinical specialists or patient experts.

#### Evidence Review Group (ERG)

An independent academic group commissioned by the National Institute for Health Research (NIHR) Health Technology Assessment Programme (HTA Programme) to assist the Appraisal Committee in reviewing the company evidence submission to the Institute.

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<sup>&</sup>lt;sup>1</sup>Non-company consultees are invited to submit statements relevant to the group they are representing.