NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Provisional Single Technology Appraisal (STA)

Liposomal cisplatin in combination with gemcitabine for previously untreated locally advanced or metastatic pancreatic cancer [ID658]

Provisional matrix of consultees and commentators

| Consultees | Commentators (no right to submit or appeal) |
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| Manufacturers/sponsors Regulon (liposomal cisplatin) Patient/carer groups Afiya Trust Black Health Agency Cancer Black Care Cancer Equality Cancer52 Equalities National Council HAWC Helen Rollason Cancer Charity Independent Cancer Patients Voice Macmillan Cancer Support Maggie's Centres Marie Curie Cancer Care Muslim Council of Britain Muslim Health Network Pancreatic Cancer Action Pancreatic Cancer UK Rarer Cancers Foundation South Asian Health Foundation Specialised Healthcare Alliance Tenovus Professional groups Association of Cancer Physicians British Geriatrics Society British Institute of Radiology British Psychosocial Oncology Society British Society of Gastroenterology Cancer Network Pharmacists Forum Cancer Research UK Pancreatic Society of Great Britain and Ireland | Allied Health Professionals Federation Board of Community Health Councils in Wales British National Formulary Care Quality Commission Department of Health, Social Services and Public Safety for Northern Ireland Healthcare Improvement Scotland Medicines and Healthcare products Regulatory Agency National Association of Primary Care National Pharmacy Association NHS Commercial Medicines Unit NHS Confederation Scottish Medicines Consortium Possible comparator manufacturers Accord Healthcare (fluorouracil, gemcitabine, irinotecan, oxaliplatin) Actavis UK (gemcitabine, irinotecan, oxaliplatin, fluorouracil) Lilly UK (gemcitabine) Medac UK (gemcitabine, irinotecan, oxaliplatin, fluorouracil,) Mylan UK (gemcitabine, irinotecan, oxaliplatin) Pfizer (irinotecan) Roche Products (capecitabine) Sandoz (fluorouracil, irinotecan) Sanofi (oxaliplatin) |
| Royal College of General Practitioners | Sun Pharmaceuticals (gemcitabine) |

National Institute for Health and Care Excellence

Provisional matrix for the proposed technology appraisal of liposomal cisplatin in combination with gemcitabine for previously untreated locally advanced or metastatic pancreatic cancer Issue date: June 2014

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Consultees Commentators (no right to submit or appeal) Royal College of Nursing Teva UK (gemcitabine, irinotecan, Royal College of Pathologists oxaliplatin) Royal College of Physicians Wockhardt UK (fluorouracil, gemcitabine, oxaliplatin) Royal College of Radiologists Zentiva (oxaliplatin) Royal Pharmaceutical Society Royal Society of Medicine Relevant research groups Society and College of Radiographers Cochrane Upper Gastrointestinal and **UK Clinical Pharmacy Association** Pancreatic Diseases Group **UK Health Forum** CORE (The Digestive Disorders) **UK Oncology Nursing Society** Foundation) Health Research Authority Others Institute of Cancer Research Department of Health MRC Clinical Trials Unit NHS England National Cancer Research Institute NHS Nottingham West CCG National Cancer Research Network NHS West Norfolk CCG National Institute of Health Research Welsh Government Pancreatic Cancer Research Fund **Evidence Review Group** Evidence Review Group tbc National Institute for Health Research Health Technology Assessment Programme **Associated Guideline Groups** National Collaborating Centre for Cancer Associated Public Health Groups Public Health England Public Health Wales

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:

Consultees

Organisations that accept an invitation to participate in the appraisal; the manufacturer(s) or sponsor(s) of the technology; national professional organisations; national patient organisations; the Department of Health and the Welsh Government and relevant NHS organisations in England.

The manufacturer/sponsor of 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-manufacturer/sponsor consultees are invited to submit a statement¹, 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: manufacturers of 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-manufacturers/sponsors 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 manufacturer/sponsor evidence submission to the Institute.

¹ Non manufacturer consultees are invited to submit statements relevant to the group they are representing.