## NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

**Review Proposal Project** 

NICE Technology Appraisal No.221; Romiplostim for the treatment of chronic immune or idiopathic thrombocytopenic purpura and No.293; Eltrombopag for treating chronic immune (idiopathic) thrombocytopenic purpura

Consultees	Commentators (no right to submit or appeal)
Company/sponsor         Amgen UK (romiplostim)         Novartis Pharmaceuticals (eltrombopag)         Patient/carer groups         Black Health Agency         Equalities National Council         ITP Support Association         Muslim Council of Britain         South Asian Health Foundation         Specialised Healthcare Alliance         Splenectomy Trust UK         Professional groups         Association of Anaesthetists         Association of Surgeons of Great Britain and Ireland         British Blood Transfusion Society         British Blood Transfusion Society         British Committee for Standards in Haematology         British Geriatrics Society         British Society for Haematology         Royal College of Anaesthetists         Royal College of Pathologists         Royal College of Physicians         Royal College of Surgeons         Royal Pharmaceutical Society         Royal Society of Medicine         UK ITP Forum	General commentators         All Wales Therapeutics and Toxicology Centre         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         Hospital Information Services - Jehovah's Witnesses         Medicines and Healthcare Products Regulatory Agency         National Association of Primary Care         National Pharmacy Association         NHS Confederation         Scottish Medicines Consortium         Welsh Health Specialised Services Committee         Comparators         Accord UK (azathioprine, mycophenolate mofetil, dapsone)         Aspen (azathioprine)         Baxter Healthcare (cyclophosphamide)         Dexcel Pharma (ciclosporin)         Hospira UK (vinblastine, vincristine)         Mylan UK (azathioprine, mycophenolate mofetil, ciclosporin, danazol)
<ul> <li>United Kingdom Clinical Pharmacy Association</li> </ul>	<ul><li>Napp Pharmaceuticals (rituximab)</li><li>Novartis (ciclosporin)</li></ul>

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Others• Department of Health and Social Care• NHS Bristol CCG• NHS England• NHS Gloucestershire CCG• Welsh Government	<ul> <li>Roche Products (mycophenolate mofetil, rituximab)</li> <li>Sandoz (rituximab, azathioprine, cyclophosphamide)</li> <li>Sanofi (danazol)</li> <li>Santen UK (ciclosporin)</li> <li>Teva Pharma (mycophenolate mofetil)</li> </ul>
	<ul> <li><u>Relevant research groups</u></li> <li>MRC Clinical Trials Unit</li> <li>National Institute for Health Research</li> </ul>
	<ul> <li><u>Associated Public Health Groups</u></li> <li>Public Health England</li> <li>Public Health Wales</li> </ul>

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

## **Consultees**

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 Social Care and the Welsh Government and relevant NHS organisations in England.

The company that markets the technology is invited to prepare a submission dossier, can respond to consultations, nominate clinical experts and has the right to appeal against the Final Appraisal Document (FAD).

All non- company consultees are invited to prepare a submission dossier respond to consultations on the draft scope, the Assessment Report and the Appraisal Consultation Document. They can nominate clinical or patient experts and have the right to appeal against the Final Appraisal Document (FAD).

## **Commentators**

Organisations that engage in the appraisal process but are not asked to prepare a submission dossier. Commentators 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; 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 the British National Formulary.

All non-company organisations can nominate clinical or patient experts to present their personal views to the Appraisal Committee.