1. **Treatment & Condition**
   Olaparib for maintenance treatment of BRCA mutation-positive advanced ovarian, fallopian tube or peritoneal cancer after response to first-line platinum-based chemotherapy.

2. **Associated appraisal body & Summary of ruling**
   NICE Technology Appraisal guidance TA598 (August 2019).
   Olaparib (Lynparza®) is recommended for use within the Cancer Drugs Fund as an option for the maintenance treatment of BRCA mutation-positive, advanced (FIGO stages 3 and 4), high-grade epithelial ovarian, fallopian tube or primary peritoneal cancer that has responded to first-line platinum-based chemotherapy in adults. It is recommended only if the conditions in the managed access agreement for olaparib are followed.

3. **Number of people in Northern Ireland expected to take up service/therapy**
   In Northern Ireland it is estimated that around 11 people per year with BRCA mutation positive, advanced (FIGO stages 3 and 4) high-grade epithelial ovarian, fallopian tube or primary peritoneal cancer that has responded to first-line platinum-based chemotherapy will be eligible for treatment with olaparib.

4. **Patient Access Scheme Availability**
   (Yes/No)
   The company (AstraZeneca) has a managed access agreement, which includes a commercial arrangement, for olaparib. The financial terms of the agreement are commercial in confidence.
   HSC Trusts will be required to claim all relevant reimbursements or discounts that form part of the managed access agreement.

5. **Infrastructure Requirements**
   Any additional infrastructure costs associated with the introduction of new cancer therapies will be dealt with as part of the routine commissioning process.

6. **Expected implementation period**
   There is no impediment to immediate implementation for new patients.
### Commissioning arrangements

This regimen will be formally commissioned by the HSCB/PHA via the Specialist Services Commissioning Team initially on a cost-per-case (CPC) basis for as long as NICE indicates that funding via the Cancer Drugs Fund is appropriate.

### Monitoring arrangements

The HSCB cost per case process will generate quarterly reports on the number of applications.

HSCB currently routinely reviews quarterly monitoring information in relation to the usage of all recurrently funded specialist cancer drugs across both the Cancer Centre and other Units.

The monitoring pro forma will be adapted to capture information in respect of this regimen and this group of patients. This monitoring report is submitted to the Specialist Services Commissioning Team for formal review and comment by the Team.

Numbers of patients who received or are receiving treatment by the HSC Board and reported to the Department of Health.

### DoH (NI) Legislative/Policy Caveats

This advice does not override or replace the individual responsibility of health professionals to make appropriate decisions in the circumstances of their individual patients, in consultation with the patient and/or guardian or carer. This would, for example, include situations where individual patients have other conditions or complications that need to be taken into account in determining whether the NICE guidance is fully appropriate in their case.

The Rural Needs Act NI 2016 has been considered and this guidance, which is purely of a technical nature, is not regarded as falling within the scope of the act.