

CervicalCheck Steering Committee

Weekly report to the Minister

14 December 2018

1. Update on support package

The provision of supports to women and families is continuing and the HSE reports that Liaison Officers are working actively to better coordinate supports and to assist in future planning for the service. The Government decision of 11 May provided that where women had been prescribed a medicine by their treating clinician, any out-of-pocket costs would be met. Arrangements are in place to reimburse, on receipt of claims, a range of costs that the women and their families incur from 11 May including travel costs, childcare costs and medical appointment costs among others. Reimbursement of retrospective costs is taking place, and an automatic review system is in place to simplify and streamline the process to ensure prompt payment of all items covered by the Government decision. To date, €1.06m has been reimbursed in respect of various health and social care costs, over €833,000 of which relates to retrospective payments. There will also be additional costs associated with the medical cards that have been issued and the meeting of certain drug costs.

A note in relation to financial support for patient advocates was submitted to the Minister for consideration in recent weeks, and the Department is now in contact with the Cervical Check Patient Support Group in relation to how best to facilitate the necessary arrangements.

2. Release of records

The HSE remains focused on responding to all requests for records as soon as possible. To aid this process external legal advisors are liaising with women and their solicitors on the release of slides. The protocol in place ensures the integrity and traceability of slides being transferred; solicitors are required to provide specific information about their chosen laboratory before slides can be released; this ensures the integrity of the slide is protected and all slides can be traced when they leave their current location. The HSE has reported that the average time to deliver slides to independent experts is 31 days.

Legal representatives for the HSE have contacted solicitors to check if they are encountering any delays and to address any issues. Arising from this exercise, a number of cases have been identified where the HSE had not been informed of the requestor's name and/or the designated lab expert to whom the slides are to be sent. As a result, the HSE is working with these solicitors to ensure that all required details are sent to the labs. The HSE provided a detailed report on the release of slides to the Public Accounts Committee last week.

3. Independent Expert Panel Review

The protocol for the International Clinical Expert Review led by the Royal College of Obstetricians and Gynaecologists (RCOG) has been published on the Department of Health website.

As of Friday 7 December, 1,702 letters, including letters to next-of-kin of women who have, sadly, died, have issued providing information in relation to consent and the consent form for participation in the review, and 1,011 women so far have consented to take part in the review (59%). A small number of letters remain to issue to next-of-kin. The transfer of slides from CervicalCheck labs for the purpose of the Expert Panel Review has encountered some external challenges which are being addressed as a matter of priority.

Validation exercise in relation to the 221 affected women

The HSE has advised it is undertaking a validation exercise on the data held on the 221 affected women. This is to ensure the National Screening Service has the most up to date information, which will be used to help with planning support needs for patients, for example.

4. Introduction of HPV as the primary method of testing

The introduction of HPV testing as the primary screening mechanism for CervicalCheck, with cytology as a reflex test, will involve a reconfiguration of the laboratory work involved. Detailed capacity planning is ongoing and introduction will be subject to a tendering process for work carried out outside the public sector. The Clinical Advisory Group established to advise on the project has made a decision in relation to the choice of HPV assay, required before tender documents can be finalised, as well as the final age range and intervals and genotyping. Dr Marc Arbyn, who was a member of the HIQA HTA panel, is also providing support and expertise to this Group. HSE officials have advised that all necessary clinical decisions which are required for drafting of procurement documents have now been made and procurement market analysis is progressing.

A pre-market engagement session took place on 27 November and submissions from those companies which attended are due to be submitted by 6 December. Work is ongoing to finalise the laboratory strategy for the HPV test, taking account of options and the futureproofing of the service. In tandem, work on development and testing of necessary ICT changes is underway and colposcopy capacity planning is being scoped out.

5. Clinical Director

The HSE has advised that the new Clinical Director has been appointed and is due to commence work in February 2019.

6. Smeartaking activity

The increased demand for smear tests has put pressure on lab capacity and the HSE has been working actively with the labs to manage this. The total number of additional GP consultations to date is over 102,000. The issue of the backlog of smear tests remains a priority concern for the HSE. Efforts to secure additional laboratory capacity to help address the backlog of smears form part of the finalisation of laboratory contract extensions.

While there is no time limit or expiry for slides in relation to cytology, there is a requirement to perform secondary HPV testing (provided where low grade abnormalities are identified in cytology, around 7-8% of all slides) within three to six months, depending on which test is used, and this is being monitored. As part of looking forward to next year, the HSE is developing a capacity plan to take account of available capacity and expected demand, with the aim of bringing the programme into stabilisation in 2019.

The provision of free GP consultations and out-of-cycle smear test is to cease, as planned, from 31 December. Letters issued to GPs on Wednesday 5 December to advise them of the current position and the CervicalCheck website has also been updated.

7. Implementation of recommendations of the Scally Inquiry

The CervicalCheck Steering Committee has overseen the development of the Implementation Plan for Dr Scally's recommendations, and agreed the plan on 4 December last. The plan has now been independently reviewed by Dr Scally, and was approved by Government on Tuesday 11 December. The Plan was subsequently published on the Department's website.

Department officials met with Dr Scally on Monday 17 December to discuss the implementation of his recommendations in more detail. It is expected that Dr Scally will provide further detailed observations formally to the Minister later in December.

The next meeting of the CervicalCheck Steering Committee is scheduled for 18 December.

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Oversight and engagement with the HSE on modules of their work as follows:

- Management of supports to patients/families
- Provision of documents to patients
- Interface with RCOG Review
- Management of laboratory capacity issues
- Introduction of HPV Screening

Management of supports to patients/families	
Significant Issues	There are no exceptional items to report in relation to Community Supports.

Provision of documents to patients	
Significant Issues	<p>Current position, issues & challenges</p> <p>The team remains focused on responding to all requests as soon as possible. External legal advisors are liaising with women and their solicitors on the release of slides. Solicitors are required to provide specific information about their chosen laboratory before slides can be released. This ensures the integrity of the slide is protected and all slides can be traced when they leave their current location. In some cases there have been delays with the laboratories due to operational pressures and we are ensuring that we continue to focus on reducing the time taken - the average time to deliver slides to the independent expert is 31 days. The HSE continues to engage directly with the laboratories to try to make sure that slides are imaged and prepared for release as quickly as possible, and to maintain ongoing pressure on the labs to improve this element of their performance. To this end, weekly operational meetings are taking place to identify, assess and deal with blockages in the process as a matter of urgency</p> <p>In relation to access to slides: some concerns were expressed in relation to significant numbers of women not getting access to slides, as a result we contacted all solicitors to establish if there were indeed people waiting on slides. As a result we have identified a significant number of cases where the HSE have not been informed of the requestors name or/and their</p>

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Interface with RCOG Review	
Project Management	<p>Project Manager in Place Assistant National Director (HSE) commenced working as the Programme Lead on the RCOG Review on Wednesday the 1st of August.</p> <p>Project Manager (HSE) commenced working as the Project Manager on Wednesday the 1st of August.</p>
Project Governance	<p>Programme Governance</p> <ul style="list-style-type: none"> Support Team holds daily meetings and teleconferences to progress deliverables, identify critical actions / timeframes, areas for escalation, and project RAIDS. Teleconference with Expert Review Panel and Dept. of Health re: project planning & matters for interagency review. <p>Next steps</p> <ul style="list-style-type: none"> Continue weekly teleconference with Dept. of Health and Expert Review Panel
Project Plan	
Workstream	Actions Progressed (w/e 7th December)
Patient Support Services	<ul style="list-style-type: none"> As of COB Fri 7th December: <ul style="list-style-type: none"> 1,702 invited to participate in Expert Panel Review, who are contactable and comprehended by the Review <i>(no change from last week)</i>. 1,011 (59%) have consented to participate in the Review <i>(increase of 87 from last week)</i>.
Information Services	<ul style="list-style-type: none"> As the number of calls to the information line have decreased to approx. 1 per day, opening hours of call centre are adjusted to 5 days per week, 9am – 6pm. As of COB Fri 7th Dec, there have been 131 calls to the information line <i>(an increase of 3 from previous week)</i>. Details as follows:

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	<table> <tr> <th>Total Calls to Information Line</th><th>#</th><th>Change from Previous Week</th></tr> <tr> <td>Total Calls (general)</td><td>131</td><td>↑ 3</td></tr> <tr> <td>Total Calls (clinical)</td><td>59</td><td>No change</td></tr> <tr> <td> Calls (clinical) resolved at initial contact</td><td>72</td><td>No change</td></tr> <tr> <td> Calls (clinical) escalated to acute hospitals, following SOP</td><td>49</td><td>No change</td></tr> <tr> <td> Calls (clinical) resolved by acute hospitals</td><td>23</td><td>No change</td></tr> </table>	Total Calls to Information Line	#	Change from Previous Week	Total Calls (general)	131	↑ 3	Total Calls (clinical)	59	No change	Calls (clinical) resolved at initial contact	72	No change	Calls (clinical) escalated to acute hospitals, following SOP	49	No change	Calls (clinical) resolved by acute hospitals	23	No change	
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Case Management System (CMS)	<ul style="list-style-type: none"> RCOG provided with colposcopy data and failsafe data from CervicalCheck ICT. 																			
Laboratory Logistics	<ul style="list-style-type: none"> CervicalCheck held extended meetings with two laboratories on 30th Nov – RCOG Programme included on agenda for discussion. 																			
Communications	No update to report																			
Acute & Community Services	<ul style="list-style-type: none"> Weekly briefing sent to HGs through Acute Operations and to Lead Colposcopists through CervicalCheck Weekly teleconference with Acute Operations Responses provided to individual clinicians who have contacted the Programme with queries on the RCOG Support Programme and the Expert Panel Review. 																			
Current Position, Significant Issues	<ul style="list-style-type: none"> The transfer of slides from CervicalCheck labs for the purpose of the Expert Panel Review is being progressed with the laboratories and issues are being worked through. 																			
Next Steps																				
Workstream	Actions Planned (w/c 10th Dec)																			

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Patient Support Services	<ul style="list-style-type: none"> • Next of kin details in relation to 9 women provided by treating hospital; letters issued December 10th. • Continue scanning and logging consent returns on Case Management System, as they are received. • Continue regular engagement with Patient Liaison Officers and other stakeholders.
Information Services	<ul style="list-style-type: none"> • Continue follow-up on calls to information line and to dedicated email address for health professionals • Employ regular quality checks to ensure capacity meets demand. Where required, identify and assign further additional clinical & administrative resources to the call centre in line with the project plan timelines.
Case Management System (CMS)	<ul style="list-style-type: none"> • As letters issue, continue to update CMS with validated data • Incorporate additional requirements for slides transfers who are also involved in legal proceedings.
Laboratory Logistics	<ul style="list-style-type: none"> • 2 Teleconferences with CC laboratories, CC Interim Programme Manager, and, RCOG Programme Lead scheduled for w/c 10th Dec to progress solutions to laboratory challenges in transferring slides.
Communications	<ul style="list-style-type: none"> • Manage feedback and queries from women/Next of Kin, clinicians, media and public as remaining cohorts of women of next of kin receive letters.
Acute & Community Services	<ul style="list-style-type: none"> • Continue weekly briefings and teleconferences with Acute Operations and CervicalCheck. • Continue developing protocol to support Expert Review Panel with relevant medical records from acute and community services where requested.

Management of Laboratory Capacity Issues	
Project Management	<p>Project Manager in Place</p> <p>Project Team Composition</p> <p>Given the nature of the issues presenting, a working group was put in place to support the project manager. This group comprises HSE procurement, HSE Legal and the CervicalCheck Operations team. This group is focused on a) extension of lab contracts and b) managing the demand due to the out of cycle smear tests.</p>
Project Plan	<p>Since April any woman who is concerned can attend her GP for a free smear and this has resulted in a testing backlog in all three labs. These delays remain a priority concern for the HSE. Because of the skilled nature of cytology screening and the difficulty in recruiting trained staff, lab turnaround time issues can be slow to resolve. Laboratories are required to capture the smear on a slide within 6 weeks. This has been</p>

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	<p>resolved with the laboratory so we do not foresee future issues. The lab working group continues to work closely with the three labs to understand and manage current capacity issues. Weekly reports are being provided and ongoing engagements to identify further actions to mitigate this risk.</p> <p>We are happy to say that we agreed continuity of supply arrangements with both existing cytology laboratories so that they continue to provide cytology services to CervicalCheck beyond the end date of the contract on Sunday 14th October. We are now concluding final contracts and operating arrangements with both Quest and Medlab pathology.</p>
Uptake of Smear Tests	<p><i>Out of cycle smears</i></p> <p>Total number of additional GP consultations to date is more than 102,000. The knock on effect of this has been 1) the estimated number of early repeat smear test to take place in the period of 1st May to 31st October is in the region of 48,088 2) a Global trend with respect to decreases in the availability of cytology staff has also exacerbated the ability of the contracted labs to increase their throughput through this time 3) as a result this has increased the reporting times to women up to 20 weeks and more in some cases. The table below sets out the increased number of GP consultations.</p> <p>In April, the Minister for Health announced that any woman of screening age with concerns about her cervical screening could avail of a free consultation, and if necessary, a free repeat smear test with any GP registered with CervicalCheck. These services will, as scheduled, cease on 31st December 2018. HSE wrote, on the 5th December 2018, to GPs registered with CervicalCheck to advise them of this process.</p>
Average Time for Processing Results	<p>The average processing time is currently 81 days.</p> <p>We are continuing to monitor the expiration of smears for low grade HPV testing. This remains a challenge for laboratories. A laboratory capacity plan is being developed for the next 6 months. This is required to stabilise the programme in order for HPV primary screening to be implemented.</p>

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Week	Total
week 1	1,946
week 2	3,739
week 3	10,683
week 4	9,335
week 5	7,946
week 6	4,213
week 7	4,797
week 8	4,198
week 9	4,110
week 10	3,174
week 11	2,907
week 12	2,737
week 13	2,823
week 14	3,061
week 15	1,949
week 16	2,422
week 17	2,397
week 18	2,871
week 19	2,534
week 20	2,427
week 21	2,338
week 22	2,130
week 23	2,571
week 24	2,098
week 25	2,126
week 26	2,026
week 27	1,922
week 28	1,737
week 29	1,642
week 30	1,727
week 31	1,422
week 32	721
Grand Total	102,729

Claim Status	Total
Fully Paid	90,934
To Pay with next Payment	10,751
Rejected	629
Duplicate	415
Grand Total	102,729



Introduction of HPV Screening

Project Management

Project Manager

In place.

Project Team Composition

- Cervical Check Clinical Director has been appointed and will commence in February 2019

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	<ul style="list-style-type: none"> The expression of interest for the appointment of a Clinical Lead Colposcopist were unsuccessful. NSS are in discussion with a pathology lead and hope to confirm an appointment for mid January. Clinical Advisory Group: international expert Dr. Marc Arbyn is supporting this group. Dr Arbyn was a member of the HIQA HTA panel and is a leading expert on HPV primary screening in Europe Still awaiting Colposcopy lead, Pathology lead.
Project Plan	
Workstream	Actions Progressed (w/e 7 th December 2018)
Clinical	<ul style="list-style-type: none"> Assessment of feedback to proposed screening pathway and eligibility framework continues.
Procurement	<ul style="list-style-type: none"> Submissions from participants by 6th December.
ICT	<ul style="list-style-type: none"> CSR amendments - testing commenced. Colposcopy IT – awaiting spec agreement, funding, and agreement of testing sites. Laboratory IT - awaiting decisions on laboratory configuration and procurement. GP IT – no specified changes yet.
Lab Configuration	<ul style="list-style-type: none"> Configuration options - work limited until the market engagement seminar is completed and current programme stabilisation w.r.t. laboratories completed. Discussion with Scottish National Cervical Screening Programme.
Communications	<ul style="list-style-type: none"> Communications lead identified for this project . To engage with team.
Resources for HPs	<ul style="list-style-type: none"> Continued work on: <ol style="list-style-type: none"> Developing training materials & the e-learning modules; and Guide for Primary Care which will be available in hardcopy and through the e-learning portal.
Hospitals (Colposcopy)	<ul style="list-style-type: none"> Colposcopy capacity planning and management of women referred to colposcopy has commenced by the women & infants programme. HG CEOs have officially been communicated to regarding this piece of work by ND NAHD

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Current Position, Significant Issues	Current position, issues & challenges <ol style="list-style-type: none"> 1) Continuing efforts to secure contract extensions with existing labs is critical to enable the programme to proceed. 2) Stabilisation of the current programme remains a huge issue. Current backlogs from the 2 private providers will impact the transition to HPV primary screening. 3) Absence of a fulltime Clinical Director for the programme is impacting a number of key activities. A Clinical Advisory Group has been set up to address these decisions. 4) Failure to recruit a colposcopy or cytopathology lead for the program and project. There were no applicants. 5) Laboratory configuration strategy being reassessed by requirements to stabilise current programme and responses to market engagement seminar. 6) Market engagement seminar – assessment of responses. 7) Communications resources (external & internal) have been identified and work now will commence on a communications plan for the transition to HPV primary screening. Urgent. 												
Immediate Activities <table border="1" data-bbox="183 868 1964 1394"> <thead> <tr> <th>Workstream</th><th>Actions Planned</th></tr> </thead> <tbody> <tr> <td>Clinical</td><td> <ul style="list-style-type: none"> Convene CAG to review clinical decisions to date and pending decisions. Circulate scope of work and arrange series of meetings with colposcopy leads. </td></tr> <tr> <td>Procurement</td><td> <ul style="list-style-type: none"> Finalise contract details with laboratories following agreement to continue. Assess outcome of market engagement seminar. </td></tr> <tr> <td>ICT</td><td> <ul style="list-style-type: none"> Continue testing of changes to Cervical Screening Register. (February 2019) Labs ICT development - awaiting nomination of labs to be contacted Colposcopy IT - Funding approval to be sought. Testing sites (colposcopy services) to be identified (Urgent). </td></tr> <tr> <td>Lab Configuration</td><td> <ul style="list-style-type: none"> Laboratory configuration strategy to be finalised looking at the options and future proofing services. </td></tr> <tr> <td>Communications</td><td> <ul style="list-style-type: none"> Assess workshop (Comms, PMO) outcomes on communications planning. </td></tr> </tbody> </table>		Workstream	Actions Planned	Clinical	<ul style="list-style-type: none"> Convene CAG to review clinical decisions to date and pending decisions. Circulate scope of work and arrange series of meetings with colposcopy leads. 	Procurement	<ul style="list-style-type: none"> Finalise contract details with laboratories following agreement to continue. Assess outcome of market engagement seminar. 	ICT	<ul style="list-style-type: none"> Continue testing of changes to Cervical Screening Register. (February 2019) Labs ICT development - awaiting nomination of labs to be contacted Colposcopy IT - Funding approval to be sought. Testing sites (colposcopy services) to be identified (Urgent). 	Lab Configuration	<ul style="list-style-type: none"> Laboratory configuration strategy to be finalised looking at the options and future proofing services. 	Communications	<ul style="list-style-type: none"> Assess workshop (Comms, PMO) outcomes on communications planning.
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Hospitals (Colposcopy)	<ul style="list-style-type: none"> Identify lead for hospitals / colposcopy requirements. Complete capacity planning exercise and schedule meetings with lead colposcopists. Secure hospital commitments for 2019. Identify colposcopy lead (clinical). 	
Update Action	<p>Update Action 04/28 Update on the possibility of appointing a clinical lead for the move to HPV testing. Update on 02/10/18-The expression of interest was extended to the 10th Oct.</p> <p>Update 19/09/18 NRS have agreed appointment date with the successful candidate for the post of clinical director.</p>	