

# Overview of the protocol

## Introduction to the Ligation Sequencing Kit V14 (SQK-LSK114) protocol

This protocol describes how to carry out sequencing of amplicon sample using the Ligation Sequencing Kit V14 (SQK-LSK114). It is recommended that a Lambda control experiment is completed first to become familiar with the technology.

For information about Kit 14 chemistry, please see the [Kit 14 sequencing and duplex basecalling](#) info sheet.

## Steps in the sequencing workflow:

### Prepare for your experiment

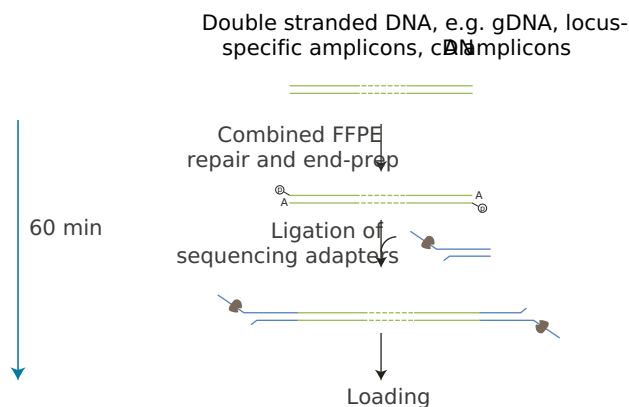
You will need to:

- Extract your DNA, and check its length, quantity and purity.**The quality checks performed during the protocol are essential in ensuring experimental success.**
- Ensure you have your sequencing kit, the correct equipment and third-party reagents
- Download the software for acquiring and analysing your data
- Check your flow cell to ensure it has enough pores for a good sequencing run

### Library preparation

The table below is an overview of the steps required in the library preparation, including timings and stopping points.

Library preparation step	Process	Time	Stop option
End-prep	Prepare the DNA ends for adapter attachment.	35 minutes	4°C overnight
Adapter ligation and clean-up	Attach sequencing adapters to the DNA ends and perform a bead clean-up.	30 minutes	4°C short-term storage or for repeated use, such as re-loading your flow cell. -80°C for single-use long-term storage. We strongly recommend sequencing your library as soon as it is adapted.
Priming and loading the flow cell	Prime the flow cell and load the prepared library for sequencing.	5 minutes	



## Sequencing and analysis

You will need to:

- Start a sequencing run using the MinKNOW software which will collect raw data and convert it into basecalled reads

### IMPORTANT

#### Compatibility of this protocol

This protocol should only be used in combination with:

- Ligation Sequencing Kit V14 (SQK-LSK114)
- Control Expansion (EXP-CTL001)
- R10.4.1 MinION Flow Cells (FLO-MIN114)
- Flow Cell Wash Kit (EXP-WSH004)
- MinION Mk1B - [MinION Mk1B IT requirements document](#)
- MinION Mk1C - [MinION Mk1C IT requirements document](#)
- GridION - [GridION IT requirements document](#)

# Equipment and consumables

## Materials

- 100-200 fmol of amplicon DNA
- Ligation Sequencing Kit V14 (SQK-LSK114)

## Consumables

- MinION and GridION Flow Cell
- NEBNext® Companion Module for Oxford Nanopore Technologies® Ligation Sequencing (NEB, E7180S or E7180L).

Alternatively, you can use the NEBNext® products below:

- NEBNext Ultra II End repair/dA-tailing Module (NEB, E7546)
- NEBNext Quick Ligation Module (NEB, E6056)
- Qubit dsDNA HS Assay Kit (Invitrogen, Q32851)
- (Optional) Bovine Serum Albumin (BSA) (50 mg/ml) (e.g. Invitrogen™ UltraPure™ BSA 50 mg/ml,

AM2616)

- Freshly prepared 80% ethanol in nuclease-free water
- Nuclease-free water (e.g. ThermoFisher, AM9937)
- 1.5 ml Eppendorf DNA LoBind tubes
- 0.2 ml thin-walled PCR tubes
- Qubit™ Assay Tubes (Invitrogen, Q32856)

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## Equipment

- MinION or GridION device
- MinION and GridION Flow Cell Light Shield
- Hula mixer (gentle rotator mixer)
- Magnetic rack, suitable for 1.5 ml Eppendorf tubes
- Microfuge
- Vortex mixer
- Thermal cycler
- P1000 pipette and tips
- P200 pipette and tips
- P100 pipette and tips
- P20 pipette and tips
- P10 pipette and tips
- P2 pipette and tips
- Ice bucket with ice
- Timer
- Qubit fluorometer (or equivalent for QC check)

**For this protocol, you will need 100-200 fmol of amplicon DNA.**

## Input DNA

### How to QC your input DNA

It is important that the input DNA meets the quantity and quality requirements. Using too little or too much DNA, or DNA of poor quality (e.g. highly fragmented or containing RNA or chemical contaminants) can affect your library preparation.

For instructions on how to perform quality control of your DNA sample, please read the [input DNA/RNA QC protocol](#).

## Chemical contaminants

Depending on how the DNA is extracted from the raw sample, certain chemical contaminants may remain in the purified DNA, which can affect library preparation efficiency and sequencing quality. Read more about contaminants on the [Contaminants page](#) of the Community.

## NEBNext® Companion Module for Oxford Nanopore Technologies® Ligation Sequencing

For customers new to nanopore sequencing, we recommend buying the [NEBNext® Companion Module](#) for Oxford Nanopore Technologies® Ligation Sequencing (catalogue number E7180S or E7180L), which contains all the NEB reagents needed for use with the Ligation Sequencing Kit.

Please note, for our amplicon protocols, NEBNext FFPE DNA Repair Mix and NEBNext FFPE DNA Repair Buffer are not required.

### Third-party reagents

We have validated and recommend the use of all the third-party reagents used in this protocol. Alternatives have not been tested by Oxford Nanopore Technologies.

For all third-party reagents, we recommend following the manufacturer's instructions to prepare the reagents for use.

### Check your flow cell

We highly recommend that you check the number of pores in your flow cell prior to starting a sequencing experiment. This should be done within three months of purchasing for MinION/GridION/PromethION or within four weeks of purchasing Flongle Flow Cells. Oxford Nanopore Technologies will replace any flow cell with fewer than the number of pores in the table below, when the result is reported within two days of performing the flow cell check, and when the storage recommendations have been followed. To do the flow cell check, please follow the instructions in the [Flow Cell Check document](#).

Flow cell	Minimum number of active pores covered by warranty
Flongle Flow Cell	50
MinION/GridION Flow Cell	800
PromethION Flow Cell	5000

#### IMPORTANT

**We strongly recommend using the Ligation Buffer (LNB) supplied in the Ligation Sequencing Kit V14 rather than the third-party ligase buffer supplied in the NEBNext Quick Ligation Module to ensure high ligation efficiency of the Ligation Adapter (LA).**

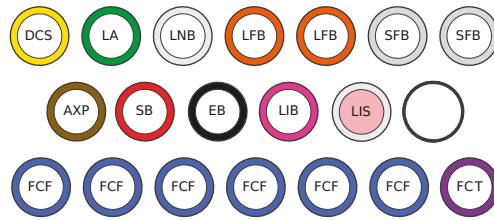
#### IMPORTANT

**Ligation Adapter (LA) included in this kit and protocol is not interchangeable with other sequencing adapters.**

### Ligation Sequencing Kit V14 (SQK-LSK114) contents

**Note:** We are in the process of reformatting our kits with single-use tubes into a bottle format.

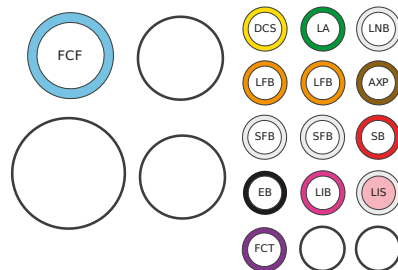
**Single-use tubes format:**



DCS : DNA Control Strand  
 LA : Ligation Adapter  
 LNB : Ligation Buffer  
 LFB : Long Fragment Buffer  
 SFB : Short Fragment Buffer  
 AXP : AMPure XP Beads

SB : Sequencing Buffer  
 EB : Elution Buffer  
 LIB : Library Beads  
 LIS : Library Solution  
 FCF : Flow Cell Flush  
 FCT : Flow Cell Tether

#### Bottle format:



DCS : DNA Control Strand  
 LA : Ligation Adapter  
 LNB : Ligation Buffer  
 LFB : Long Fragment Buffer  
 SFB : Short Fragment Buffer  
 AXP : AMPure XP Beads

SB : Sequencing Buffer  
 EB : Elution Buffer  
 LIB : Library Beads  
 LIS : Library Solution  
 FCF : Flow Cell Flush  
 FCT : Flow Cell Tether

**Note:** This Product Contains AMPure XP Reagent Manufactured by Beckman Coulter, Inc. and can be stored at -20°C with the kit without detriment to reagent stability.

**Note:** The DNA Control Sample (DCS) is a 3.6 kb standard amplicon mapping the 3' end of the Lambda genome.

## End-prep

~35 minutes

#### Materials

- 100-200 fmol amplicon DNA
- DNA Control Sample (DCS)
- AMPure XP Beads (AXP)

#### Consumables

- NEBNext® Ultra II End Repair / dA-tailing Module (NEB, E7546)
- Qubit dsDNA HS Assay Kit (Invitrogen, Q32851)
- Freshly prepared 80% ethanol in nuclease-free water
- Nuclease-free water (e.g. ThermoFisher, AM9937)
- 0.2 ml thin-walled PCR tubes
- 1.5 ml Eppendorf DNA LoBind tubes
- Qubit™ Assay Tubes (Invitrogen, Q32856)

## Equipment

- P1000 pipette and tips
- P100 pipette and tips
- P10 pipette and tips
- Thermal cycler
- Microfuge
- Hula mixer (gentle rotator mixer)
- Magnetic rack
- Ice bucket with ice
- Qubit fluorometer (or equivalent for QC check)

Check your flow cell.

We recommend performing a flow cell check before starting your library prep to ensure you have a flow cell with enough pores for a good sequencing run.

See the [flow cell check instructions](#) in the MinKNOW protocol for more information.

**1 Thaw DNA Control Sample (DCS) at room temperature, spin down, mix by pipetting, and place on ice.**

**2 Prepare the NEBNext Ultra II End Repair / dA-tailing Module reagents in accordance with manufacturer's instructions, and place on ice:**

For optimal performance, NEB recommend the following:

1. Thaw all reagents on ice.
2. Flick and/or invert the reagent tubes to ensure they are well mixed.

**Note:** Do not vortex the Ultra II End Prep Enzyme Mix.

3. Always spin down tubes before opening for the first time each day.
4. The Ultra II End Prep Buffer may have a little precipitate. Allow the mixture to come to room temperature and pipette the buffer up and down several times to break up the precipitate, followed by vortexing the tube for 30 seconds to solubilise any precipitate.

### IMPORTANT

**Do not vortex the NEBNext Ultra II End Prep Enzyme Mix.**

### IMPORTANT

**It is important that the NEBNext Ultra II End Prep Reaction Buffer is mixed well by vortexing.**

Check for any visible precipitate; vortexing for at least 30 seconds may be required to solubilise all precipitate.

**3 Prepare the DNA in nuclease-free water:**

- Transfer 100-200 fmol of amplicon DNA into a 1.5 ml Eppendorf DNA LoBind tube
- Adjust the volume to 49 µl with nuclease-free water
- Mix thoroughly by pipetting up and down, or by flicking the tube
- Spin down briefly in a microfuge

**4 In a 0.2 ml thin-walled PCR tube, mix the following:**

Between each addition, pipette mix 10-20 times.

Reagent	Volume
DNA CS	1 µl
DNA	49 µl
Ultra II End-prep Reaction Buffer	7 µl
Ultra II End-prep Enzyme Mix	3 µl
<b>Total</b>	<b>60 µl</b>

**5 Thoroughly mix the reaction by gently pipetting and briefly spinning down.**

**6 Using a thermal cycler, incubate at 20°C for 5 minutes and 65°C for 5 minutes.**

**7 Resuspend the AMPure XP Beads (AXP) by vortexing.**

**8 Transfer the DNA sample to a clean 1.5 ml Eppendorf DNA LoBind tube.**

**9 Add 60 µl of resuspended the AMPure XP Beads (AXP) to the end-prep reaction and mix by flicking the tube.**

**10 Incubate on a Hula mixer (rotator mixer) for 5 minutes at room temperature.**

**11 Prepare 500 µl of fresh 80% ethanol in nuclease-free water.**

**12 Spin down the sample and pellet on a magnet until supernatant is clear and colourless. Keep the tube on the magnet, and pipette off the supernatant.**

**13 Keep the tube on the magnet and wash the beads with 200 µl of freshly prepared 80% ethanol without disturbing the pellet. Remove the ethanol using a pipette and discard.**

**14 Repeat the previous step.**

**15 Spin down and place the tube back on the magnet. Pipette off any residual ethanol. Allow to dry for ~30 seconds, but do not dry the pellet to the point of cracking.**

**16 Remove the tube from the magnetic rack and resuspend the pellet in 61 µl nuclease-free water. Incubate for 2 minutes at room temperature.**

**17 Pellet the beads on a magnet until the eluate is clear and colourless, for at least 1 minute.**

**18 Remove and retain 61 µl of eluate into a clean 1.5 ml Eppendorf DNA LoBind tube.**

Quantify 1 µl of eluted sample using a Qubit fluorometer.

#### END OF STEP

**Take forward the repaired and end-prepped DNA into the adapter ligation step. However, at this point it is also possible to store the sample at 4°C overnight.**

## Adapter ligation and clean-up

~30 minutes

### Materials

- Ligation Adapter (LA)
- Ligation Buffer (LNB) from the Ligation Sequencing Kit
- Long Fragment Buffer (LFB)
- Short Fragment Buffer (SFB)
- AMPure XP Beads (AXP)
- Elution Buffer (EB) from the Oxford Nanopore sequencing kit

### Consumables

- Qubit dsDNA HS Assay Kit (Invitrogen, Q32851)
- NEBNext Quick Ligation Module (NEB, E6056)
- 1.5 ml Eppendorf DNA LoBind tubes
- Qubit™ Assay Tubes (Invitrogen, Q32856)

### Equipment

- Magnetic rack
- Microfuge
- Vortex mixer
- P1000 pipette and tips
- P100 pipette and tips
- P20 pipette and tips
- P10 pipette and tips
- Qubit fluorometer (or equivalent for QC check)



#### IMPORTANT

Ligation Adapter (LA) included in this kit and protocol is not interchangeable with other sequencing adapters.

#### IMPORTANT

Although the recommended third-party ligase is supplied with its own buffer, the ligation efficiency of the Ligation Adapter (LA) is higher when using the Ligation Buffer (LNB) supplied in the Ligation Sequencing Kit.

- 1 Spin down the Ligation Adapter (LA) and Quick T4 Ligase, and place on ice.
- 2 Thaw Ligation Buffer (LNB) at room temperature, spin down and mix by pipetting. Due to viscosity, vortexing this buffer is ineffective. Place on ice immediately after thawing and mixing.
- 3 Thaw the Elution Buffer (EB) at room temperature and mix by vortexing. Then spin down and place on ice.

#### IMPORTANT

Depending on the wash buffer (LFB or SFB) used, the clean-up step after adapter ligation is designed to either enrich for DNA fragments of >3 kb, or purify all fragments equally.

- To enrich for DNA fragments of 3 kb or longer, use Long Fragment Buffer (LFB)
- To retain DNA fragments of all sizes, use Short Fragment Buffer (SFB)

- 4 Thaw either Long Fragment Buffer (LFB) or Short Fragment Buffer (SFB) at room temperature and mix by vortexing. Then spin down and place on ice.
- 5 In a 1.5 ml Eppendorf DNA LoBind tube, mix in the following order:  
Between each addition, pipette mix 10-20 times.

Reagent	Volume
DNA sample from the previous step	60 µl
Ligation Buffer (LNB)	25 µl
NEBNext Quick T4 DNA Ligase	10 µl
Ligation Adapter (LA)	5 µl
<b>Total</b>	<b>100 µl</b>

- 6 Thoroughly mix the reaction by gently pipetting and briefly spinning down.
- 7 Incubate the reaction for 10 minutes at room temperature.

- 8 Resuspend the AMPure XP Beads (AXP) by vortexing.
  - 9 Add 40 µl of resuspended AMPure XP Beads (AXP) to the reaction and mix by flicking the tube.
  - 10 Incubate on a Hula mixer (rotator mixer) for 5 minutes at room temperature.
  - 11 Spin down the sample and pellet on a magnet. Keep the tube on the magnet, and pipette off the supernatant when clear and colourless.
  - 12 Wash the beads by adding either 250 µl Long Fragment Buffer (LFB) or 250 µl Short Fragment Buffer (SFB). Flick the beads to resuspend, spin down, then return the tube to the magnetic rack and allow the beads to pellet. Remove the supernatant using a pipette and discard.
  - 13 Repeat the previous step.
  - 14 Spin down and place the tube back on the magnet. Pipette off any residual supernatant. Allow to dry for ~30 seconds, but do not dry the pellet to the point of cracking.
  - 15 Remove the tube from the magnetic rack and resuspend the pellet in 15 µl Elution Buffer (EB). Spin down and incubate for 10 minutes at room temperature. For high molecular weight DNA, incubating at 37°C can improve the recovery of long fragments.
  - 16 Pellet the beads on a magnet until the eluate is clear and colourless, for at least 1 minute.
  - 17 Remove and retain 15 µl of eluate containing the DNA library into a clean 1.5 ml Eppendorf DNA LoBind tube.  
Dispose of the pelleted beads
- Quantify 1 µl of eluted sample using a Qubit fluorometer.
- 18 Prepare 35-50 fmol of your final library to 12 µl with Elution Buffer (EB).

#### IMPORTANT

We recommend loading 35-50 fmol of this final prepared library onto the R10.4.1 flow cell.

This is to ensure high pore occupancy of >95% is reached. How to calculate pore occupancy can be found [here](#).

#### END OF STEP

The prepared library is used for loading into the flow cell. Store the library on ice or at 4°C until ready to load.

#### TIP

##### Library storage recommendations

We recommend storing libraries in Eppendorf DNA LoBind tubes at **4°C for short-term** storage or repeated use, for example, re-loading flow cells between washes.

For single use and **long-term storage** of more than 3 months, we recommend storing libraries at **-80°C** in Eppendorf DNA LoBind tubes.

# Priming and loading the MinION and GridION Flow Cell

~5 minutes

## Materials

- Flow Cell Flush (FCF)
- Flow Cell Tether (FCT)
- Library Solution (LIS)
- Library Beads (LIB)
- Sequencing Buffer (SB)

## Consumables

- MinION and GridION Flow Cell
- 1.5 ml Eppendorf DNA LoBind tubes
- Nuclease-free water (e.g. ThermoFisher, AM9937)
- (Optional) Bovine Serum Albumin (BSA) (50 mg/ml) (e.g. Invitrogen™ UltraPure™ BSA 50 mg/ml, AM2616)

## Equipment

- MinION or GridION device
- MinION and GridION Flow Cell Light Shield
- P1000 pipette and tips
- P100 pipette and tips
- P20 pipette and tips
- P10 pipette and tips

#### IMPORTANT

Please note, this kit is only compatible with R10.4.1 flow cells (FLO-MIN114).

#### TIP

##### Priming and loading a flow cell

We recommend all new users watch the [Priming and loading your flow cell](#) video before your first run.

- 1 Thaw the Sequencing Buffer (SB), Library Beads (LIB) or Library Solution (LIS, if using), Flow Cell Tether (FCT) and Flow Cell Flush (FCF) at room temperature before mixing by vortexing. Then spin down and store on ice.

IMPORTANT

For optimal sequencing performance and improved output on MinION R10.4.1 flow cells (FLO-MIN114), we recommend adding Bovine Serum Albumin (BSA) to the flow cell priming mix at a final concentration of 0.2 mg/ml.

**Note:** We do not recommend using any other albumin type (e.g. recombinant human serum albumin).

- 2 To prepare the flow cell priming mix with BSA, combine Flow Cell Flush (FCF) and Flow Cell Tether (FCT), as directed below. Mix by pipetting at room temperature.

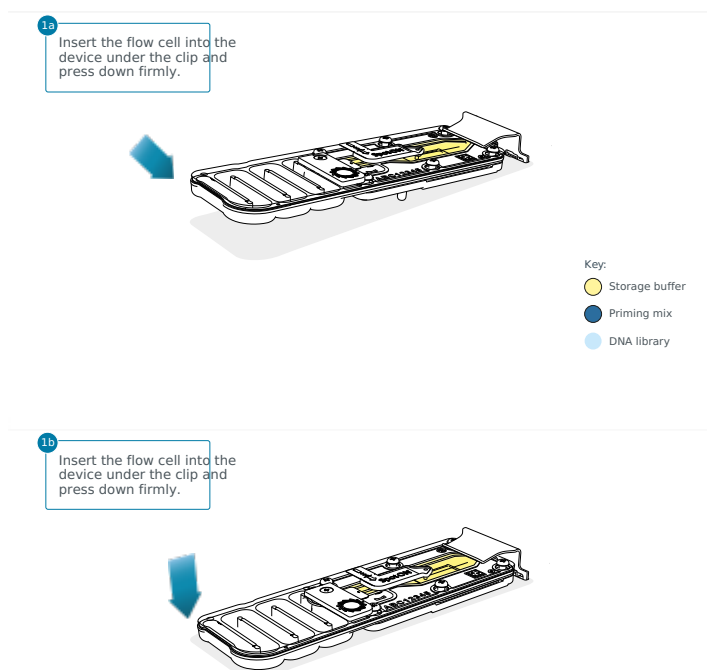
**Note:** We are in the process of reformatting our kits with single-use tubes into a bottle format. Please follow the instructions for your kit format.

**Single-use tubes format:**  
Add 5 µl Bovine Serum Albumin (BSA) at 50 mg/ml and 30 µl Flow Cell Tether (FCT) directly to a tube of Flow Cell Flush (FCF).

**Bottle format:**  
In a suitable tube for the number of flow cells, combine the following reagents:

Reagent	Volume per flow cell
Flow Cell Flush (FCF)	1,170 µl
Bovine Serum Albumin (BSA) at 50 mg/ml	5 µl
Flow Cell Tether (FCT)	30 µl
<b>Total volume</b>	<b>1,205 µl</b>

- 3 Open the MinION or GridION device lid and slide the flow cell under the clip. Press down firmly on the flow cell to ensure correct thermal and electrical contact.**



**Optional action**

Complete a flow cell check to assess the number of pores available before loading the library.

This step can be omitted if the flow cell has been checked previously.

See the [flow cell check instructions](#) in the MinKNOW protocol for more information.

- 4 Slide the flow cell priming port cover clockwise to open the priming port.**

### IMPORTANT

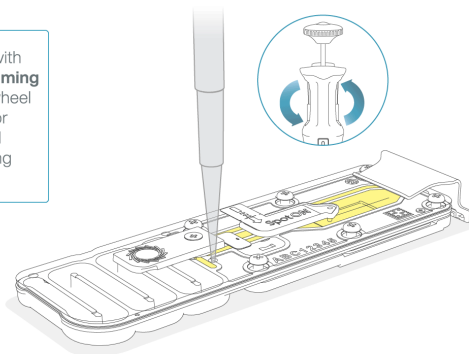
Take care when drawing back buffer from the flow cell. Do not remove more than 20-30  $\mu\text{l}$ , and make sure that the array of pores are covered by buffer at all times. Introducing air bubbles into the array can irreversibly damage pores.

#### 5 After opening the priming port, check for a small air bubble under the cover. Draw back a small volume to remove any bubbles:

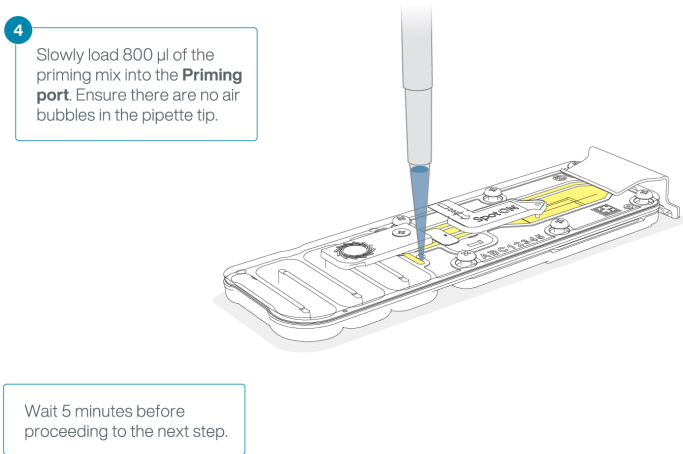
1. Set a P1000 pipette to 200  $\mu\text{l}$
2. Insert the tip into the priming port
3. Turn the wheel until the dial shows 220-230  $\mu\text{l}$ , to draw back 20-30  $\mu\text{l}$ , or until you can see a small volume of buffer entering the pipette tip

**Note:** Visually check that there is continuous buffer from the priming port across the sensor array.

3 Insert a P1000 pipette with an empty tip into the **Priming port**. Turn the pipette wheel to draw back 20-30  $\mu\text{l}$  or until you can see a small volume of buffer entering the pipette tip.



6 Load 800 µl of the priming mix into the flow cell via the priming port, avoiding the introduction of air bubbles. Wait for five minutes. During this time, prepare the library for loading by following the steps below.



7 Thoroughly mix the contents of the Library Beads (LIB) by pipetting.

**IMPORTANT**

The Library Beads (LIB) tube contains a suspension of beads. These beads settle very quickly. It is vital that they are mixed immediately before use.

We recommend using the Library Beads (LIB) for most sequencing experiments. However, the Library Solution (LIS) is available for more viscous libraries.

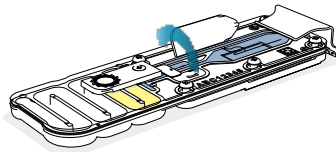
8 In a new 1.5 ml Eppendorf DNA LoBind tube, prepare the library for loading as follows:

Reagent	Volume per flow cell
Sequencing Buffer (SB)	37.5 µl
Library Beads (LIB) mixed immediately before use, or Library Solution (LIS), if using	25.5 µl
DNA library	12 µl
<b>Total</b>	<b>75 µl</b>

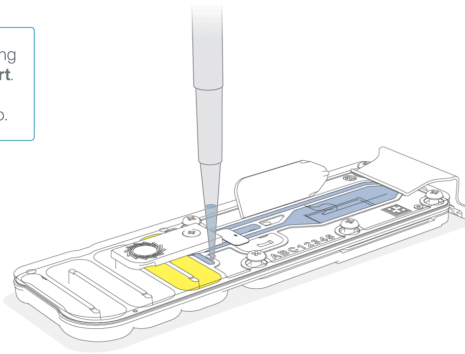
## 9 Complete the flow cell priming:

1. Gently lift the SpotON sample port cover to make the SpotON sample port accessible.
2. Load **200 µl** of the priming mix into the flow cell priming port (**not** the SpotON sample port), avoiding the introduction of air bubbles.

5 Gently flip open the SpotON sample port cover.



6 Load 200 µl of the priming mix into the **Priming Port**. Ensure there are no air bubbles in the pipette tip.

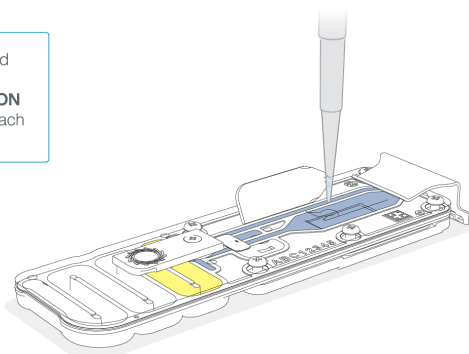


## 10 Mix the prepared library gently by pipetting up and down just prior to loading.



**11 Add 75 µl of the prepared library to the flow cell via the SpotON sample port in a dropwise fashion. Ensure each drop flows into the port before adding the next.**

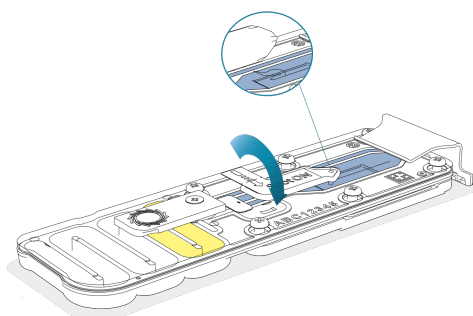
**7** Pipette mix the prepared library and load 75 µl dropwise into the **SpotON** sample port, ensuring each drop flows into the port.



**12 Gently replace the SpotON sample port cover, making sure the bung enters the SpotON port and close the priming port.**

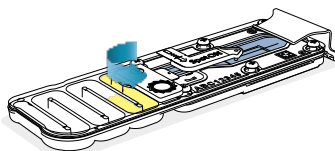
8

Gently replace the **SpotON** sample port cover.



9

Gently close the **Priming port**.



#### **IMPORTANT**

**Install the light shield on your flow cell as soon as library has been loaded for optimal sequencing output.**

We recommend leaving the light shield on the flow cell when library is loaded, including during any washing and reloading steps. The shield can be removed when the library has been removed from the flow cell.

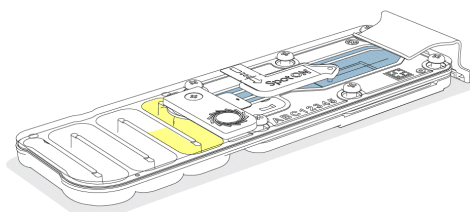
### 13 Place the light shield onto the flow cell, as follows:

1. Carefully place the leading edge of the light shield against the clip.

**Note:** Do not force the light shield underneath the clip.

2. Gently lower the light shield onto the flow cell. The light shield should sit around the SpotON cover, covering the entire top section of the flow cell.

10 Carefully align the **light shield** against the clip and lower onto the flow cell.



#### CAUTION

The MinION Flow Cell Light Shield is not secured to the flow cell and careful handling is required after installation.

#### END OF STEP

Close the device lid and set up a sequencing run on MinKNOW.

## Data acquisition and basecalling

### How to start sequencing

Once you have loaded your flow cell, the sequencing run can be started on MinKNOW, our sequencing software that controls the device, data acquisition and real-time basecalling. For more detailed information on setting up and using MinKNOW, please see the [MinKNOW protocol](#).

MinKNOW can be used and set up to sequence in multiple ways:

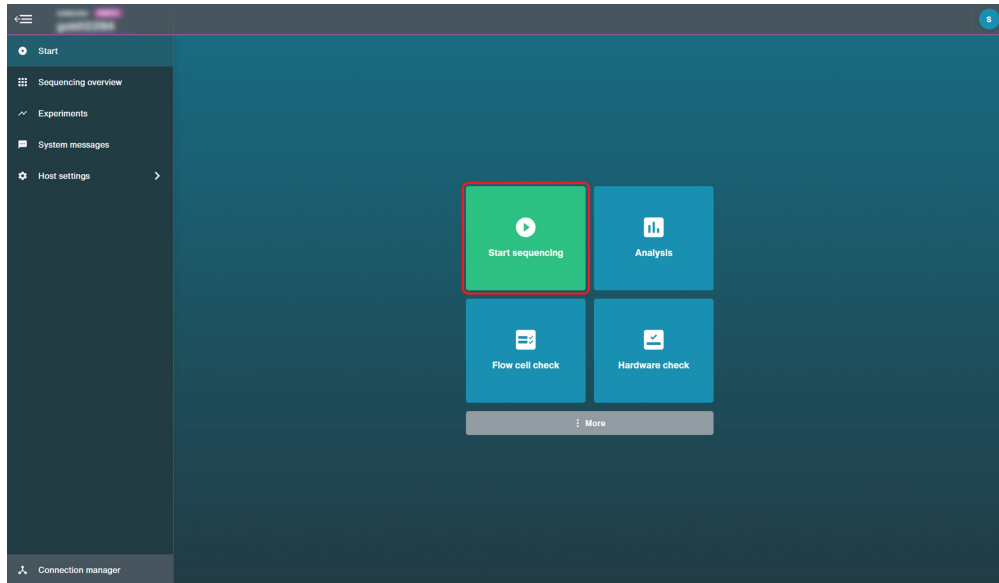
- On a computer either directly or remotely connected to a sequencing device.
- Directly on a GridION, MinION Mk1C or PromethION 24/48 sequencing device.

For more information on using MinKNOW on a sequencing device, please see the device user manuals:

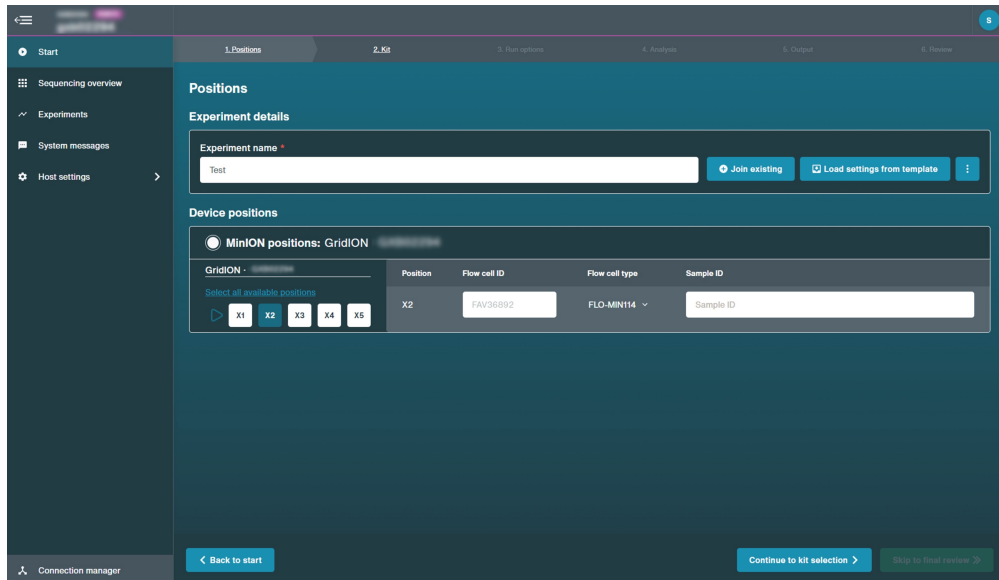
- [MinION Mk1B user manual](#)
- [MinION Mk1C user manual](#)
- [GridION user manual](#)

To start a sequencing run on MinKNOW:

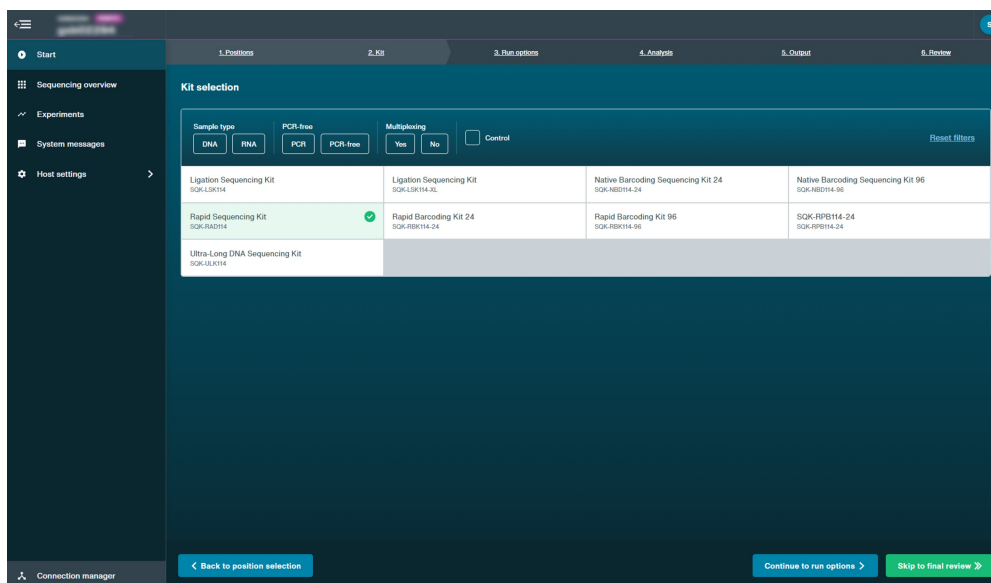
1. Navigate to the start page and click **Start sequencing**.



2. Fill in your experiment details, such as name and flow cell position and sample ID.



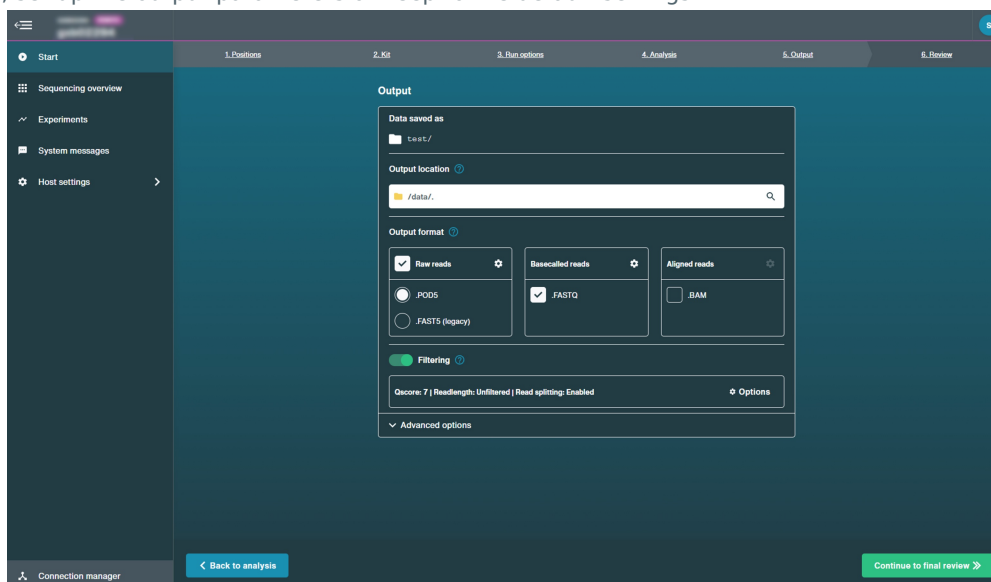
3. Select the sequencing kit used in the library preparation on the Kit page.



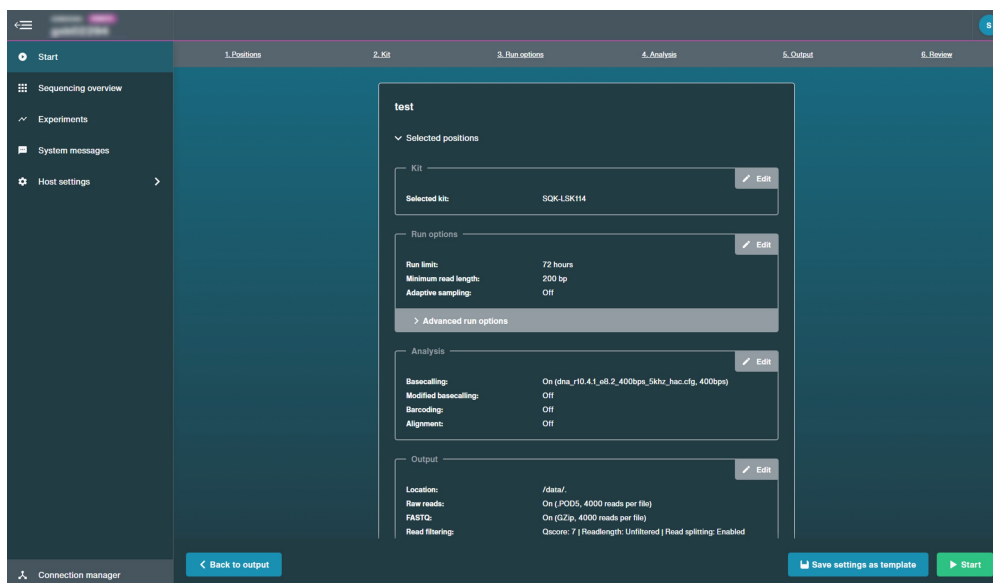
4. Configure the sequencing parameters for your sequencing run or keep to the default settings on the Run options and Analysis tabs.

**Note:** If basecalling was turned off when a sequencing run was set up, basecalling can be performed post-run on MinKNOW. For more information, please see the [MinKNOW protocol](#).

5. On the Output page, set up the output parameters or keep to the default settings.



6. Click **Start** on the Review page to start the sequencing run.



## Duplex basecalling

Kit 14 chemistry has improved duplex basecalling which requires rebasecalling on [Dorado](#) after simplex basecalling has been performed on MinKNOW.

For detailed information on setting up your sequencing run, for both simplex and duplex basecalling, please see the [Kit 14 sequencing and duplex basecalling](#) info sheet.

**Note:** When running Dorado, we recommend stopping other basecalling for the best performance by maximising memory available to Dorado. This can be stopped and restarted when Dorado has finished via the GUI on MinKNOW.

## Data analysis after sequencing

After sequencing has completed on MinKNOW, the flow cell can be reused or returned, as outlined in the Flow cell reuse and returns section.

After sequencing and basecalling, the data can be analysed. For further information about options for basecalling and post-basecalling analysis, please refer to the [Data Analysis](#) document.

In the Downstream analysis section, we outline further options for analysing your data.

# Flow cell reuse and returns

## Materials

- Flow Cell Wash Kit (EXP-WSH004)

- 1 **After your sequencing experiment is complete, if you would like to reuse the flow cell, please follow the Flow Cell Wash Kit protocol and store the washed flow cell at 2-8°C.**

The [Flow Cell Wash Kit protocol](#) is available on the Nanopore Community.

#### TIP

**We recommend you to wash the flow cell as soon as possible after you stop the run. However, if this is not possible, leave the flow cell on the device and wash it the next day.**

- 2 **Alternatively, follow the returns procedure to flush out the flow cell ready to send back to Oxford Nanopore.**

Instructions for returning flow cells can be found [here](#).

**Note:** All flow cells must be flushed with deionised water before returning the product.

#### IMPORTANT

**If you encounter issues or have questions about your sequencing experiment, please refer to the Troubleshooting Guide that can be found in the online version of this protocol.**

## Downstream analysis

### Post-basecalling analysis

There are several options for further analysing your basecalled data:

#### 1. EPI2ME workflows

For in-depth data analysis, Oxford Nanopore Technologies offers a range of bioinformatics tutorials and workflows available in EPI2ME Labs, which are available in the [EPI2ME Labs](#) section of the Community. The platform provides a vehicle where workflows deposited in GitHub by our Research and Applications teams can be showcased with descriptive texts, functional bioinformatics code and example data.

#### 2. Research analysis tools

Oxford Nanopore Technologies' Research division has created a number of analysis tools, that are available in the Oxford Nanopore [GitHub repository](#). The tools are aimed at advanced users, and contain instructions for how to install and run the software. They are provided as-is, with minimal support.

#### 3. Community-developed analysis tools

If a data analysis method for your research question is not provided in any of the resources above, please refer to the [Bioinformatics](#) section of the Resource centre. Numerous members of the Nanopore Community have developed their own tools and pipelines for analysing nanopore sequencing data, most of which are available on GitHub. Please be aware that these tools are not supported by Oxford Nanopore Technologies, and are not guaranteed to be compatible with the latest chemistry/software configuration.

# Issues during DNA/RNA extraction and library preparation

Below is a list of the most commonly encountered issues, with some suggested causes and solutions.

We also have an FAQ section available on the [Nanopore Community Support](#) section.

If you have tried our suggested solutions and the issue still persists, please contact Technical Support via email ([support@nanoporetech.com](mailto:support@nanoporetech.com)) or via [LiveChat](#) in the Nanopore Community.

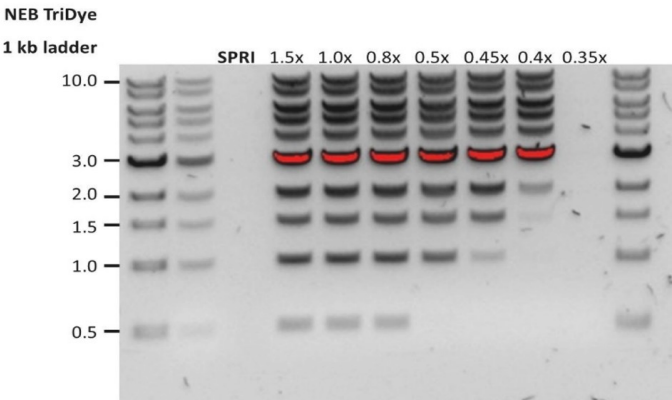
## Low sample quality

Observation	Possible cause	Comments and actions
<b>Low DNA purity (Nanodrop reading for DNA OD 260/280 is &lt;1.8 and OD 260/230 is &lt;2.0-2.2)</b>	The DNA extraction method does not provide the required purity	The effects of contaminants are shown in the <a href="#">Contaminants</a> document. Please try an alternative <a href="#">extraction method</a> that does not result in contaminant carryover.  Consider performing an additional SPRI clean-up step.
<b>Low RNA integrity (RNA integrity number &lt;9.5 RIN, or the rRNA band is shown as a smear on the gel)</b>	The RNA degraded during extraction	Try a different <a href="#">RNA extraction method</a> . For more info on RIN, please see the <a href="#">RNA Integrity Number</a> document. Further information can be found in the <a href="#">DNA/RNA Handling</a> page.
<b>RNA has a shorter than expected fragment length</b>	The RNA degraded during extraction	Try a different <a href="#">RNA extraction method</a> . For more info on RIN, please see the <a href="#">RNA Integrity Number</a> document. Further information can be found in the <a href="#">DNA/RNA Handling</a> page.  We recommend working in an RNase-free environment, and to keep your lab equipment RNase-free when working with RNA.

## Low DNA recovery after AMPure bead clean-up

Observation	Possible cause	Comments and actions
<b>Low recovery</b>	DNA loss due to a lower than intended AMPure beads-to-sample ratio	1. AMPure beads settle quickly, so ensure they are well resuspended before adding them to the sample.  2. When the AMPure beads-to-sample ratio is lower than 0.4:1, DNA fragments of any size will be lost during the clean-up.



Observation	Possible cause	Comments and actions
<b>Low recovery</b>	DNA fragments are shorter than expected	<p>The lower the AMPure beads-to-sample ratio, the more stringent the selection against short fragments. Please always determine the input DNA length on an agarose gel (or other gel electrophoresis methods) and then calculate the appropriate amount of AMPure beads to use.</p> 
<b>Low recovery after end-prep</b>	The wash step used ethanol <70%	DNA will be eluted from the beads when using ethanol <70%. Make sure to use the correct percentage.

## Issues during the sequencing run

Below is a list of the most commonly encountered issues, with some suggested causes and solutions.

We also have an FAQ section available on the [Nanopore Community Support](#) section.

If you have tried our suggested solutions and the issue still persists, please contact Technical Support via email ([support@nanoporetech.com](mailto:support@nanoporetech.com)) or via [LiveChat](#) in the Nanopore Community.

### Fewer pores at the start of sequencing than after Flow Cell Check

Observation	Possible cause	Comments and actions
<b>MinKNOW reported a lower number of pores at the start of sequencing than the number reported by the Flow Cell Check</b>	An air bubble was introduced into the nanopore array	After the Flow Cell Check it is essential to remove any air bubbles near the priming port before priming the flow cell. If not removed, the air bubble can travel to the nanopore array and irreversibly damage the nanopores that have been exposed to air. The best practice to prevent this from happening is demonstrated in <a href="#">this video</a> .

Observation	Possible cause	Comments and actions
<b>MinKNOW reported a lower number of pores at the start of sequencing than the number reported by the Flow Cell Check</b>	The flow cell is not correctly inserted into the device	Stop the sequencing run, remove the flow cell from the sequencing device and insert it again, checking that the flow cell is firmly seated in the device and that it has reached the target temperature. If applicable, try a different position on the device (GridION/PromethION).
<b>MinKNOW reported a lower number of pores at the start of sequencing than the number reported by the Flow Cell Check</b>	Contaminations in the library damaged or blocked the pores	The pore count during the Flow Cell Check is performed using the QC DNA molecules present in the flow cell storage buffer. At the start of sequencing, the library itself is used to estimate the number of active pores. Because of this, variability of about 10% in the number of pores is expected. A significantly lower pore count reported at the start of sequencing can be due to contaminants in the library that have damaged the membranes or blocked the pores. Alternative DNA/RNA extraction or purification methods may be needed to improve the purity of the input material. The effects of contaminants are shown in the <a href="#">Contaminants Know-how piece</a> . Please try an alternative <a href="#">extraction method</a> that does not result in contaminant carryover.

#### MinKNOW script failed

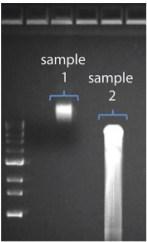
Observation	Possible cause	Comments and actions
<b>MinKNOW shows "Script failed"</b>		Restart the computer and then restart MinKNOW. If the issue persists, please collect the <a href="#">MinKNOW log files</a> and contact Technical Support. If you do not have another sequencing device available, we recommend storing the flow cell and the loaded library at 4°C and contact Technical Support for further storage guidance.

#### Pore occupancy below 40%

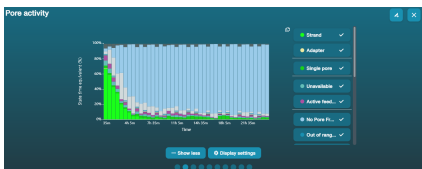
Observation	Possible cause	Comments and actions
<b>Pore occupancy &lt;40%</b>	Not enough library was loaded on the flow cell	Ensure the correct volume and concentration as stated on the appropriate protocol for your sequencing library is loaded onto the flow cell. Please quantify the library before loading and calculate fmols using tools like the <a href="#">Promega Biomath Calculator</a> , choosing "dsDNA: µg to fmol"
<b>Pore occupancy close to 0</b>	The Ligation Sequencing Kit was used, and sequencing adapters did not ligate to the DNA	Make sure to use the NEBNext Quick Ligation Module (E6056) and Oxford Nanopore Technologies Ligation Buffer (LNB, provided in the SQK-LSK110 kit) at the sequencing adapter ligation step, and use the correct amount of each reagent. A Lambda control library can be prepared to test the integrity of the third-party reagents.
<b>Pore occupancy close to 0</b>	The Ligation Sequencing Kit was used, and ethanol was used instead of LFB or SFB at the wash step after sequencing adapter ligation	Ethanol can denature the motor protein on the sequencing adapters. Make sure the LFB or SFB buffer was used after ligation of sequencing adapters.

Observation	Possible cause	Comments and actions
<b>Pore occupancy close to 0</b>	No tether on the flow cell	Tethers are adding during flow cell priming (FLT tube for Kit 9, 10, 11, and FCT for Kit 14). Make sure FLT/FCT was added to the buffer (FB for Kit 9, 10, 11, and FCF for Kit 14) before priming.

### Shorter than expected read length

Observation	Possible cause	Comments and actions
<b>Shorter than expected read length</b>	Unwanted fragmentation of DNA sample	<p>Read length reflects input DNA fragment length. Input DNA can be fragmented during extraction and library prep.</p> <ol style="list-style-type: none"> <li>1. Please review the <a href="#">Extraction Methods</a> in the Nanopore Community for best practice for extraction.</li> <li>2. Visualise the input DNA fragment length distribution on an agarose gel before proceeding to the library prep.</li> </ol>  <p>In the image above, Sample 1 is of high molecular weight, whereas Sample 2 has been fragmented.</p> <ol style="list-style-type: none"> <li>3. During library prep, avoid pipetting and vortexing when mixing reagents. Flicking or inverting the tube is sufficient.</li> </ol>

### Large proportion of unavailable pores

Observation	Possible cause	Comments and actions
<p><b>Large proportion of unavailable pores (shown as blue in the channels panel and pore activity plot)</b></p>  <p>The pore activity plot above shows an increasing proportion of "unavailable" pores over time.</p>	Contaminants are present in the sample	<p>Some contaminants can be cleared from the pores by the unblocking function built into MinKNOW. If this is successful, the pore status will change to "sequencing pore". If the portion of unavailable pores stays large or increases:</p> <ol style="list-style-type: none"> <li>1. A <a href="#">nuclease flush using the Flow Cell Wash Kit (EXP-WSH004)</a> can be performed, or</li> <li>2. Run several cycles of PCR to try and dilute any contaminants that may be causing problems.</li> </ol>

### Large proportion of inactive pores

Observation	Possible cause	Comments and actions
<b>Large proportion of inactive/unavailable pores (shown as light blue in the channels panel and pore activity plot. Pores or membranes are irreversibly damaged)</b>	Air bubbles have been introduced into the flow cell	Air bubbles introduced through flow cell priming and library loading can irreversibly damage the pores. Watch the <a href="#">Priming and loading your flow cell</a> video for best practice
<b>Large proportion of inactive/unavailable pores</b>	Certain compounds co-purified with DNA	Known compounds, include polysaccharides, typically associate with plant genomic DNA.  1. Please refer to the <a href="#">Plant leaf DNA extraction method</a> . 2. Clean-up using the QIAGEN PowerClean Pro kit. 3. Perform a whole genome amplification with the original gDNA sample using the QIAGEN REPLI-g kit.
<b>Large proportion of inactive/unavailable pores</b>	Contaminants are present in the sample	The effects of contaminants are shown in the <a href="#">Contaminants</a> Know-how piece. Please try an alternative extraction method that does not result in contaminant carryover.

#### Reduction in sequencing speed and q-score later into the run

Observation	Possible cause	Comments and actions
<b>Reduction in sequencing speed and q-score later into the run</b>	For Kit 9 chemistry (e.g. SQK-LSK109), fast fuel consumption is typically seen when the flow cell is overloaded with library (please see the appropriate protocol for your DNA library to see the recommendation).	Add more fuel to the flow cell by following the instructions in the <a href="#">MinKNOW protocol</a> . In future experiments, load lower amounts of library to the flow cell.

#### Temperature fluctuation

Observation	Possible cause	Comments and actions
<b>Temperature fluctuation</b>	The flow cell has lost contact with the device	Check that there is a heat pad covering the metal plate on the back of the flow cell. Re-insert the flow cell and press it down to make sure the connector pins are firmly in contact with the device. If the problem persists, please contact Technical Services.

#### Failed to reach target temperature

Observation	Possible cause	Comments and actions
<b>MinKNOW shows "Failed to reach target temperature"</b>	The instrument was placed in a location that is colder than normal room temperature, or a location with poor ventilation (which leads to the flow cells overheating)	MinKNOW has a default timeframe for the flow cell to reach the target temperature. Once the timeframe is exceeded, an error message will appear and the sequencing experiment will continue. However, sequencing at an incorrect temperature may lead to a decrease in throughput and lower q-scores. Please adjust the location of the sequencing device to ensure that it is placed at room temperature with good ventilation, then re-start the process in MinKNOW. Please refer to <a href="#">this FAQ</a> for more information on MinION Mk 1B temperature control.

