



# MarLIN

## Marine Information Network

Information on the species and habitats around the coasts and sea of the British Isles

# ***Laminaria hyperborea* forest with a faunal cushion (sponges and polyclinids) and foliose red seaweeds on very exposed upper infralittoral rock**

MarLIN – Marine Life Information Network  
Marine Evidence-based Sensitivity Assessment (MarESA) Review

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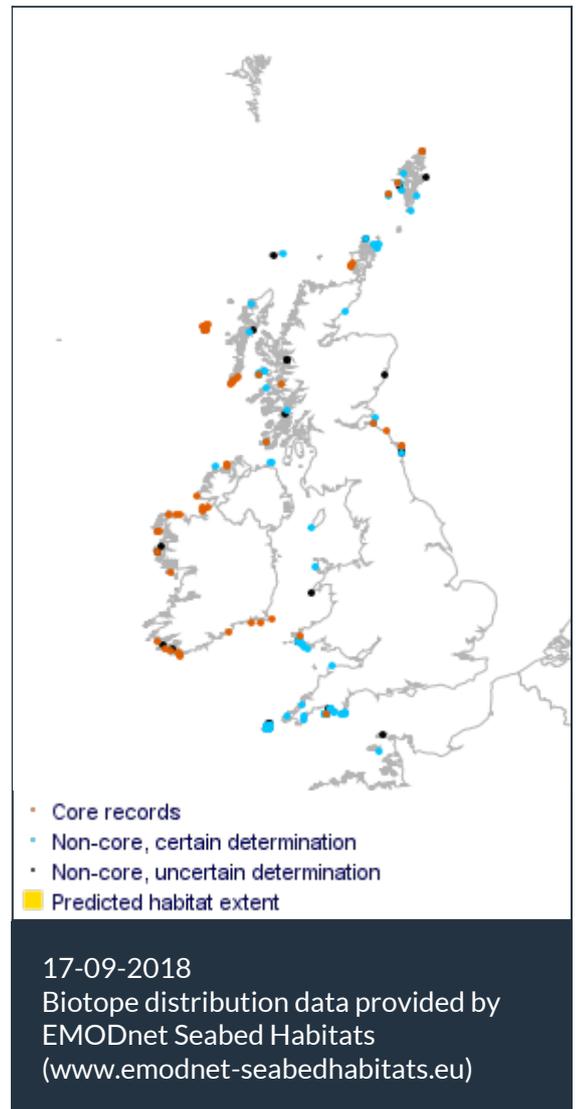
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*Laminaria hyperborea* forest with a faunal cushion (sponges and polyclinids) and foliose red seaweeds on very exposed upper infralittoral rock.

Photographer: Bernard Picton

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Researched by Thomas Stamp and Dr Keith Hiscock

Refereed by This information is not refereed.

## Summary

### ☰ UK and Ireland classification

EUNIS 2008 A3.113

*Laminaria hyperborea* forest with a faunal cushion (sponges and polyclinids) and foliose red seaweeds on very exposed infralittoral rock

JNCC 2015 IR.HIR.KFaR.LhypFa

*Laminaria hyperborea* forest with a faunal cushion (sponges and polyclinids) and foliose red seaweeds on very exposed upper infralittoral rock

JNCC 2004 IR.HIR.KFaR.LhypFa

*Laminaria hyperborea* forest with a faunal cushion (sponges and polyclinids) and foliose red seaweeds on very exposed upper infralittoral rock

1997 Biotope IR.EIR.KFaR.LhypFa

*Laminaria hyperborea* forest with a faunal cushion (sponges and polyclinids) and foliose red seaweeds on very exposed infralittoral rock

### 🔍 Description

Very exposed and exposed, but wave-surfed, upper infralittoral bedrock and massive boulders characterised by a dense forest of the kelp *Laminaria hyperborea* with a high diversity of seaweeds and invertebrates. The shallowest kelp plants are often short or stunted, while deeper plants are taller with heavily epiphytised stipes with foliose red seaweeds such as *Delesseria sanguinea*, *Cryptopleura ramosa* or *Plocamium cartilagineum* or even the brown seaweed *Dictyota dichotoma*. Also found on the stipes or on the rock below the canopy are red seaweeds including *Phycodrys rubens*, *Kallymenia reniformis*, *Callophyllis laciniata*, *Caryophyllia smithii*, and *Corallina officinalis*, while encrusting coralline algae can cover any bare patches of rock. At some sites the red seaweeds can be virtually mono-specific, while at other sites show considerable variation containing a dense mixed turf of a large variety of species. The red seaweed *Odonthalia dentata* can be present in the north. The faunal and floral under-storey is generally rich in species due, in part, to the relatively low urchin-grazing pressure in such shallow exposed conditions. The faunal composition of this biotope varies markedly between sites, but commonly occurring are the soft coral *Alcyonium digitatum* and the anthozoans *Sagartia elegans* and *Corynactis viridis*. Sponges form a prominent part of the community with variable amounts of the sponges *Halichondria panicea* and *Pachymatisma johnstonia* and several other species. The crab *Cancer pagurus* and the starfish *Asterias rubens* are normally present in small numbers foraging beneath the canopy, while the sea urchins *Echinus esculentus* and *Urticina felina* graze on the seaweeds. The hydroid *Obelia geniculata*, the ascidian *Botryllus schlosseri* and the bryozoan *Membranipora membranacea* compete for space on the kelp, whereas the bryozoan *Electra pilosa* also can be found on foliose red seaweeds.

This kelp forest most commonly occurs beneath a zone of *Alaria esculenta* and *Mytilus edulis* (Ala.Myt) and may contain small patches of *Alaria esculenta*. As the force of the wave-surge diminishes with increased depth, density of the faunal turf reduces and the kelp forest or park changes to one characterized by kelp and dense red seaweeds (LhypR.Ft or LhypR.Pk). In some areas of Shetland and St Kilda the lower infralittoral zone is characterised by a park of the kelp *Saccharina latissima* and/or *Saccorhiza polyschides* (SlatSac). Where the *Laminaria hyperborea* forest continues to depths of 15 m or greater it may give way to a zone of dense foliose red algae (FoR or For.Dic). (Information from Connor *et al.*, 2004; JNCC, 2015).

## ↓ Depth range

0-5 m, 5-10 m, 10-20 m

## 🏛️ Additional information

-

## ✓ Listed By

- none -

## 🔗 Further information sources

Search on:



## Habitat review

### 🔄 Ecology

#### Ecological and functional relationships

This is an extremely dynamic biotope with the main rock cover species that occupy it competing for space and significant seasonal changes occurring. In shallow depths, suspension feeding animal species may out-compete algae and dominate even in well-lit areas. Also, sea urchins are often absent from shallow (say, less than 10m) depths due to strong wave action thus allowing a much more lush growth of algae than would be the case if grazing was occurring. Kelps are major primary producers and up to 90 percent of kelp production enters the detrital food web so that kelp is probably a major contributor of organic carbon to surrounding communities (Birkett *et al.* 1998b). Kelp fronds, stipes and holdfasts provide substrata for distinct communities of species, some of which are found only or especially on kelp plants. Hiscock & Mitchell (1980) list 15 species of algae associated with kelp stipes in the UK. The stipes also support epifaunal bryozoa and hydroids (Norton *et al.* 1977). Holdfasts support a diverse fauna that represents a sample of the surrounding mobile fauna and crevice dwelling organisms, e.g., polychaetes, small crabs, gastropods, bivalves, and amphipods. Jones (1971) lists 53 macrofaunal invertebrates in holdfasts and Moore (1973) reports 389 species from holdfasts collected in the north east coast of Britain. An account of holdfast fauna is given by Hayward (1988). Where sea-urchins occur, they graze the undercanopy and understory algae, including juvenile kelp sporophytes, together with epiphytes and epifauna on the lower reaches of the laminarian stipe. Sea urchin grazing may maintain the patchy and species rich understory epiflora/fauna by preventing a small number of species from becoming dominant. Vost (1983) examined the effect of removing grazing *Echinus esculentus* and found that after 6-10 months the patchiness of the understory algae had decreased and the species richness and biomass of epilithic species increased. *Echinus esculentus* grazing probably controls the lower limit of *Laminaria hyperborea* distribution in some location, e.g. in the Isle of Man (Jones & Kain 1967; Kain *et al.* 1974; Kain 1979). Other ecological relationships are:

- Epiphytes and understory algae are grazed by a variety of amphipods, isopods and gastropods, e.g. *Littorina* spp., *Acmaea* spp., *Aplysia* and rissoid gastropods (Birkett *et al.* 1988b).
- Lobsters (*Homarus gammarus*), crabs and some fish species (e.g. the wolffish *Anarchicas lupus*) and perhaps otters are known to consume gastropod and echinoderm grazers.
- Kelp communities and the interaction between kelp, urchins and predators has been studied in Nova Scotia, Norway, southern California and the UK (Mann 1982; Kain 1979; Sivertsen 1997; Vadas & Elner 1992; Elner & Vadas, 1990; Schiel & Foster, 1986).
- Birkett *et al.* (1998b) suggest that juveniles of animals present in kelp beds as adults probably use the habitat as a nursery and unknown numbers of species are likely to use the habitat during their life cycle. Rinde *et al.* (1992 cited in Birkett *et al.* 1998b) state that Norwegian kelp beds are nurseries for gadoid species.
- The composition of the holdfast fauna has been shown to vary with turbidity (natural and anthropogenic in origin), between kelp species (due to holdfast architecture and volume), and with location around the coast of the British Isles (Moore 1973a&b; Moore 1978; Edwards 1980; Sheppard *et al.* 1980). Moore (1973a&b) identified groups of species that were found in most cases, or restricted to either turbid or clear waters. Moore (1978) noted that species diversity or amphipods decreased with increasing turbidity, partly due to the increased dominance of a few species. Edwards (1980) noted that holdfast fauna in south-west Ireland were numerically dominated by suspension feeders with decreasing

numbers of omnivores and carnivores respectively. Edwards (1980) noted that holdfasts were dominated by *Spirobranchus triqueter* in the most turbid sites, although these were not as turbid as sites examined by Moore (1973a&b). Along the North Sea coast species number and diversity increased with increased clarity, however where heavy metals were a factor species number and diversity decreased with increasing heavy metal pollution. They were able to distinguish groups of species characteristic of all sites, or clear or turbid sites. Along the west coast both heavy metals and turbidity were important. Where turbidity and heavy metals increased suspension feeders increased in abundance while other trophic groups decreased. However, along the south coast longitude was the most important factor, and they suggested that natural variation in temperature, salinity and water flow were responsible for variation between holdfast communities (Sheppard *et al.* 1980). Moore (1985) also demonstrated that the amphipod fauna varied with water flow rate (resulting from wave action and currents); for example sites of increased exposure were dominated by *Amphithoe rubricata*, *Lembos websteri* and *Jassa falcata* whereas *Gitana sarsi*, *Dexamine thea* and *Corophium bonnellii* flourish in wave sheltered environments.

- A few meiofaunal species may burrow into kelp tissue, e.g. the nematode *Monohystera disjuncta* (Birkett *et al.* 1998b).
- The understory flora varies with location, depth, exposure, hydrographic regime, turbidity and siltation and may be sparse or species rich. Birkett *et al.* (1998b; Appendix 5) list 52 common kelp biotope understory algae in the UK including characterizing species such as *Delesseria sanguinea*, *Dictyota dichotoma*, *Phycodrys rubens*, *Cryptopleura ramosa*, *Plocamium cartilagineum*, and *Callophyllis laciniata*.
- The benthic fauna varies with depth, exposure, location and substratum, however, no species are specific to kelp forest. Norton *et al.* (1977) demonstrate the zonation of 22 epibenthic species. However, many species, both fixed and mobile, are present and probably under recorded (Birkett *et al.* 1998b).

### Seasonal and longer term change

The species present in the biotope are believed to be mainly present throughout the year. However, many algae will show a seasonal change from recruitment of ephemeral species and re-growth of perennial species in the spring, through growth of epibiota on the algae in summer and degeneration of fronds in many species in the autumn and winter. New blades of *Laminaria hyperborea* grow in winter between the meristem and the old blade, which is shed in early spring or summer together with associated species growing on its surface. Larger and older kelp plants become liable to removal by wave action and storms due to their size and weakening by grazers such as *Patella pellucida*. There is therefore likely to be a reduced abundance of kelps following the winter. Loss of older plants results in more light reaching the understory, temporarily permitting growth of algae including *Laminaria hyperborea* sporelings. Areas of kelp may become denuded of macroalgae at intervals and the substrata dominated by encrusting corallines. These areas are often associated with an increase in urchin numbers forming 'fronts' of small and large urchins that remove large quantities of algae including the kelps themselves forming 'urchin barrens'. Sea urchin grazing is an important factor in kelp beds and, as part of the biotope, the following suggested factors affecting sea urchin populations are presented.

- Several predators have been suggested as controlling sea urchin populations e.g. sea otters, lobsters, crabs or wolffish, however the evidence is equivocal (Birkett *et al.* 1998b; Elner & Vadas 1990; Mann 1982).
- Evidence suggests that sea urchin recruitment is sporadic and may be enhanced by low temperatures (Birkett *et al.* 1998b).

- Sea urchin recruitment is also enhanced by the presence of 'urchin barrens' presumably due to the lack of suspension feeders that would otherwise consume their larvae (Lang & Mann 1978).
- Sea urchin diseases, such as 'bald-urchin' disease, encouraged by high water temperatures drastically reduce the urchin population (Lobban & Harrison, 1997). However, although parasitic infections are found in *Echinus esculentus*, no evidence of sea urchin disease has been found in the UK.
- Sivertsen (1997) examined grazing of west and north Norwegian coast *Laminaria hyperborea* beds by *Strongylocentrotus droebachiensis* and *Echinus esculentus*. He concluded that seven environmental factors contributed to the distribution of kelp beds and 'barrens': depth gradient, latitude, time of sampling, nematode infection (in *Strongylocentrotus droebachiensis*), wave exposure, coastal gradient and substratum.

The factors controlling sea urchin populations and 'urchin barrens' in kelp beds is poorly understood, especially in the UK. However, it is likely that the local urchin population is controlled by a number of factors that vary between sites and biotopes; including predators, competition for food with other grazers, variation in sea urchin recruitment, and parasitic infection or disease. Periodic storms are likely to remove older and weaker plants creating patches cleared of kelp and increasing the local turbidity. While cleared patches may encourage growth of sporelings or gametophyte maturation, they may also enhance sea urchin recruitment.

### Habitat structure and complexity

Kelp beds are diverse species rich habitats and over 1,800 species have been recorded in the UK kelp biotopes (Birkett *et al.* 1998b). Kelp forest provides a variety of habitats and refugia in a similar way to terrestrial forests. Kelps also reduce current flow producing a sheltered microclimate. In kelp forest (e.g. EIR.LhypR.Ft) the kelp density produces a canopy which excludes up to 90 % of incident light allowing many deeper water, shade tolerant algae, mainly reds, to invade. In deeper water, as light intensity decreases, the kelp density decreases forming a kelp park (Norton *et al.* 1977). Kelp beds are patchy and dynamic with areas devoid of kelp (due to storms, wave surge or grazing) in the process of expansion or recolonization in different stages of succession. Species diversity changes with depth, between forest & park, with exposure, substratum and turbidity (Birkett *et al.* 1998b; Erwin *et al.* 1990; Norton *et al.* 1977).

1. Nekton: wrasse and pollock have been observed associated especially with kelp forests and epibenthic predatory or herbivorous fish are also found, e.g. blennies, gobies and wolffish (*Anarhichas lupus*).

### Productivity

Kelps are the major primary producers in UK marine coastal waters producing nearly 75 percent of the net carbon fixed annually on the shoreline of the coastal euphotic zone (Birkett *et al.* 1998b). Kelp plants produce 2.7 times their standing biomass per year. Kelp detritus, as broken plant tissue, particles and dissolved organic material supports soft bottom communities outside the kelp bed itself. The kelps reduce ambient levels of nutrients, although this may not be significant in exposed sites, but increase levels of particulate and dissolved organic matter within the bed.

### Recruitment processes

Recruitment processes of key characteristic or dominant species are described here. *Laminaria hyperborea* produces vast numbers of spores, however they need to settle and form gametophytes within about 1 mm of each to ensure fertilisation and therefore may suffer from dilution effects

over distance. Gametophytes can survive darkness and develop in the low light levels under the canopy. However, young sporelings develop slowly in low light. Loss of older plants provides the opportunity to develop into adult plants. Recruitment in *Echinus esculentus* is sporadic or annual depending on location and may benefit from the presence of 'urchin barrens'. *Patella pellucida* is an annual species, larvae settling in the lower eulittoral and juveniles migrating to kelp, via several algal species, as they grow. (Please view individual key information reviews for details). Epifaunal larvae probably contribute to the plankton of the kelp bed and many are lost to the suspension feeding epifauna. Kelp beds also provide nurseries for larvae and fish species (see above). Recruitment of epiphytes and epiflora are dependant on dispersal and settlement of algal spores and survival of early post-settlement stages. Norton (1992) suggests that spore dispersal is primarily dependant on currents and eddies. Settlement of algal spores is partly dependant on their motility (if any) and adhesive properties together with preferences for topography (surface roughness), the chemical nature of the substratum and water movement (Norton 1992; Fletcher & Callow 1992). Vadas *et al.* 1992 suggested that survival of early post settlement stages is dependant on grazing, the algal canopy and turf effects together with desiccation and water motion, and they further suggest that recruitment is likely to be episodic, variable and to suffer from high mortality of early stages. Kain (1975) examined recolonization of artificially cleared areas in a *Laminaria hyperborea* forest in Port Erin, Isle of Man. Cleared concrete blocks were colonized by *Saccorhiza polychides*, *Alaria esculenta*, *Desmarestia* spp., *Laminaria hyperborea*, *Laminaria digitata*, *Saccharina latissima* (studied as *Laminaria saccharina*) and un-specified Rhodophyceae at 0.8m. *Saccorhiza polychides* dominated within 8 months but had virtually disappeared with 77 weeks to be replaced by laminarians, including *Alaria esculenta*. After about 2.5 years, *Laminaria hyperborea* standing crop, together with an understorey of red algae (Rhodophyceae), was similar to that of virgin forest. Rhodophyceae were present throughout the succession increasing from 0.04 to 1.5 percent of the biomass within the first 4 years. Colonizing species varied with time of year, for example blocks cleared in August 1969 were colonized by primarily *Saccharina latissima* and subsequent colonization by *Laminaria hyperborea* and other laminarians was faster than blocks colonized by *Saccorhiza polychides*; within 1 year the block was occupied by laminarians and Rhodophyceae only. Succession was similar at 4.4m, and *Laminaria hyperborea* dominated within about 3 years. Blocks cleared in August 1969 at 4.4m were not colonized by *Saccorhiza polychides* but were dominated by Rhodophyceae after 41 weeks, e.g. *Delesseria sanguinea* and *Cryptopleura ramosa*. Kain (1975) cleared one group of blocks at two monthly intervals and noted that Phaeophyceae were dominant colonists in spring, Chlorophyceae (solely *Ulva lactuca*) in summer and Rhodophyceae were most important in autumn and winter. Animal species are likely to recruit mainly from the plankton although some species such as polychinid tunicates may have larva that swim for only 2-3 hours (Berril 1950) or no larval stage (amphipods). Little is known about the reproductive biology and dispersal of some species but information from clearance experiments (see 'Time for community to reach maturity') suggests that sponges may be slow to settle.

### Time for community to reach maturity

Experimental clearance experiments in the Isle of Man (Kain 1975; Kain, 1979) showed that *Laminaria hyperborea* out-competed other opportunistic species (e.g. *Alaria esculenta*, *Saccorhiza polychides* and *Desmarestia* spp.) and returned to near control levels of biomass within 3 years at 0.8 m but that recovery was slower at 4.4m (see above). Kain (1979) noted that grazing would slow recovery as few sporophytes survived after 1 year in the presence of *Echinus esculentus*. However, the presence of other kelps and *Desmarestia* spp. (the latter is distasteful to grazers due to presence of sulphuric acids in its tissue) may act as refugia from grazing for developing *Laminaria hyperborea* juveniles that eventually out-compete the other species. Studies of the effects of

harvesting in Norway (Svendsen 1972 cited in Birkett *et al.* 1998b) showed that kelp biomass returned 3-4 years after harvesting, although the plants were small (about 1m) and the age class was shifted towards younger plants. Sivertsen (1991 cited in Birkett *et al.* 1998b), showed that kelp populations stabilise about 4-5 years after harvesting. Re-growth partly due to growth of viable juveniles remaining after harvesting. Current advice suggests that kelp forest should be left 7-10 years for kelp and non-kelp species to recover (Birkett *et al.* 1998b). Detailed studies in Norway by Rinde *et al.* (1992 cited in Birkett *et al.* 1998b) examined recovery of non-kelp species. The epiphyte community in control areas about 10 years old was richer and more extensive than on replacement plants in harvested areas. Of the epifauna, *Halichondria* sp. were only found on 10 year old plants and tunicates on plants 6 years post harvesting. Holdfast fauna was more abundant richer in 10 year old plants in control areas than younger plants in previously harvested area. Older plants have larger holdfasts. Shrimp, lobster, hermit crabs, *Echinus esculentus* and *Strongylocentrotus droebachiensis* were associated with holdfasts in control areas but absent from harvested areas. Control areas had a more diverse benthic macroflora and macrofauna. Dredged areas exhibited growth of opportunistic kelps e.g. *Alaria esculenta* and also *Desmarestia* spp. while the bottom was covered by coralline algae between young *Laminaria hyperborea* after 3 years. Control areas had a more diverse bottom community. Overall his results suggest that full biological recovery, or maturation, may take at least 10 years.

### Additional information

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## Preferences & Distribution

### Habitat preferences

Depth Range	0-5 m, 5-10 m, 10-20 m
<a href="#">Water clarity preferences</a>	
Limiting Nutrients	No information found
Salinity preferences	Full (30-40 psu)
Physiographic preferences	Open coast
Biological zone preferences	Upper infralittoral
Substratum/habitat preferences	Bedrock, Large to very large boulders
Tidal strength preferences	Moderately Strong 1 to 3 knots (0.5-1.5 m/sec.), Strong 3 to 6 knots (1.5-3 m/sec.), Weak < 1 knot (<0.5 m/sec.)
Wave exposure preferences	Exposed, Extremely exposed, Very exposed
Other preferences	High wave exposure

### Additional Information

## Species composition

### Species found especially in this biotope

### Rare or scarce species associated with this biotope

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### Additional information

The biotope is defined by conspicuous species but is probably rich in a wide range of other mobile animal species especially amongst the foliose algae, in kelp holdfasts and attached to the kelp stipes and fronds.

## Sensitivity review

### Sensitivity characteristics of the habitat and relevant characteristic species

At high densities, *Laminaria hyperborea* forms a canopy over infralittoral rock. Beneath the canopy an understory community grows, typically defined by a red seaweed turf although faunal species dominate in tide swept and/or wave surged conditions. Grazing by the urchins; *Echinus esculentus* and *Paracentrotus lividus* can also define the biotope and reduce the biomass of *Laminaria hyperborea* and understory flora. The abundance of *Laminaria hyperborea* is determined by light availability, which decreases with an increase in water depth. Therefore, depth and water clarity determines the density of *Laminaria* and hence the distribution of kelp forest (high density kelp) and park (low density kelp) sub-biotopes.

Kelp biotopes are a major source of primary productivity, and support magnified secondary productivity within North Atlantic coastal waters (Smale *et al.*, 2013, Brodie *et al.*, 2014). In Scotland, alone kelp biotopes are estimated to cover 8000km<sup>2</sup> (Walker, 1953), and account for ca 45% of primary production in UK coastal waters (Smale *et al.*, 2013). Therefore kelp biotopes, of which *Laminaria hyperborea* is dominant within UK subtidal rocky reefs (Birkett *et al.*, 1998), make a substantial contribution to coastal primary production in the UK (Smale *et al.*, 2013). *Laminaria hyperborea* is grazed directly by species such as *Patella pellucida*, however approximately 80% of primary production is consumed as detritus or dissolved organic material (Krumhansl, 2012) which is both retained within and transported out of the parent kelp forest, providing valuable nutrition to potentially low productivity habitats such as sandy beaches (Smale *et al.*, 2013).

*Laminaria hyperborea* also acts as an ecosystem engineer (Jones *et al.*, 1996; Smale *et al.*, 2013) by altering; light levels (Sjötun *et al.*, 2006), physical disturbance (Connell, 2003), sedimentation rates (Eckman *et al.*, 1989) and water flow (Smale *et al.*, 2013), profoundly altering the physical environment for fauna and flora in close proximity. *Laminaria hyperborea* biotopes increase the three-dimensional complexity of unvegetated rock (Norderhaug, 2004, Norderhaug *et al.*, 2007, Norderhaug & Christie, 2011, Gorman *et al.*, 2012; Smale *et al.*, 2013) and support high local diversity, abundance and biomass of epi/benthic species (Smale *et al.*, 2013), and serve as a nursery ground for a number of commercial important species, e.g. Gadidae (The taxonomic family that contains many commercially important marine fish species, including the Atlantic Cod and Pollack) (Rinde *et al.*, 1992).

In undertaking this assessment of sensitivity, an account is taken of knowledge of the biology of all characterizing species/taxa in the biotope. For this sensitivity assessment *Laminaria hyperborea* is the primary focus of research, however, it is recognized that the understory community, typically red seaweeds, also define the biotope. Examples of important species groups are mentioned where appropriate.

### Resilience and recovery rates of habitat

A number of review and experimental publications have assessed the recovery of *Laminaria hyperborea* kelp beds and the associated community. If environmental conditions are favourable *Laminaria hyperborea* can recover following disturbance events reaching comparable plant densities and size to pristine *Laminaria hyperborea* beds within 2-6 years (Kain, 1979; Birkett *et al.*, 1998b; Christie *et al.*, 1998). Holdfast communities may recover in 6 years (Birkett *et al.*, 1998b). Full epiphytic community and stipe habitat complexity regeneration require over 6 years (possibly 10 years). These recovery rates were based on discrete kelp harvesting events. Recurrent

disturbance occurring frequently within 2-6 years of the initial disturbance is likely to lengthen recovery time (Birkett *et al.*, 1998b, Burrows *et al.*, 2014). Kain (1975) cleared sublittoral blocks of *Laminaria hyperborea* at different times of the year for several years. The first colonizers and succession community differed between blocks and at what time of year the blocks were cleared, however within 2 years of clearance the blocks were dominated by *Laminaria hyperborea* (Fletcher *et al.*, 2006).

In south Norway, *Laminaria hyperborea* forests are harvested, which results in large scale removal of the canopy-forming kelps. Cristie *et al.*, (1998) found that in south Norwegian *Laminaria hyperborea* beds a pool of small (<25cm) understory *Laminaria hyperborea* plants persist beneath the kelp canopy for several years. The understory *Laminaria hyperborea* sporophytes had fully re-established the canopy at a height of 1m within 2-6 years after kelp harvesting. Within 1 year following harvesting, and each successive year thereafter, a pool of *Laminaria hyperborea* recruits had re-established within the understory beneath the kelp canopy. Cristie *et al.*, (1998) suggested that *Laminaria hyperborea* bed re-establishment from understory recruits (see above) inhibits the colonization of other kelps species and furthers the dominance of *Laminaria hyperborea* within suitable habitats, stating that *Laminaria hyperborea* habitats are relatively resilient to disturbance events.

*Laminaria hyperborea* has a heteromorphic life strategy, A vast number of zoospores (mobile asexual spores) are released into the water column between October-April (Kain & Jones, 1964). Zoospores settle onto rock substrata and develop into dioecious gametophytes (Kain, 1979) which, following fertilization, develop into sporophytes and mature within 1-6 years (Kain, 1979; Fredriksen *et al.*, 1995; Christie *et al.*, 1998). *Laminaria hyperborea* zoospores have a recorded dispersal range of ~200m (Fredriksen *et al.*, 1995). However zoospore dispersal is greatly influenced by water movements, and zoospore density and the rate of successful fertilization decreases exponentially with distance from the parental source (Fredriksen *et al.*, 1995). Hence, recruitment following disturbance can be influenced by the proximity of mature kelp beds producing viable zoospores to the disturbed area. (Kain, 1979, Fredriksen *et al.*, 1995).

*Laminaria hyperborea* biotopes are partially reliant on low (or no) populations of sea urchins, primarily the species; *Echinus esculentus*, *Paracentrotus lividus* and *Strongylocentrotus droebachiensis*, which graze directly on macroalgae, epiphytes and the understory community. Multiple authors (Steneck *et al.*, 2002; Steneck *et al.*, 2004; Rinde & Sjøtun, 2005; Norderhaug & Christie, 2009; Smale *et al.*, 2013) have reported dense aggregations of sea urchins to be a principal threat to *Laminaria hyperborea* biotopes of the North Atlantic. Intense urchin grazing creates expansive areas known as “urchin barrens”, in which a shift can occur from *Laminaria hyperborea* dominated biotopes to those characterized by coralline encrusting algae, with a resultant reduction in biodiversity (Lienaas & Christie, 1996; Steneck *et al.*, 2002; Norderhaug & Christie, 2009). Continued intensive urchin grazing pressure on *Laminaria hyperborea* biotopes can inhibit the *Laminaria hyperborea* recruitment (Sjøtun *et al.*, 2006) and cause urchin barrens to persist for decades (Cristie *et al.*, 1998; Steneck *et al.*, 2004; Rinde & Sjøtun, 2005). The mechanisms that control sea urchin aggregations are poorly understood but have been attributed to anthropogenic pressure on top down urchin predators (e.g. cod or lobsters). While these theories are largely unproven, a number of studies have shown that removal of urchins from grazed areas coincides with kelp re-colonization (Lienaas & Christie, 1996; Norderhaug & Christie, 2009). Lienaar & Christie, (1996) removed *Strongylocentrotus droebachiensis* from “urchin barrens” and observed a succession effect, in which the substratum was initially colonized by filamentous macroalgae and *Saccharina latissima*. However, after 2-4 years *Laminaria hyperborea* dominated the community.

Reports of large-scale urchin barrens within the North East Atlantic are generally limited to regions of the North Norwegian and Russian Coast (Rinde & Sjøtun, 2005, Nourderhaug & Christie, 2009). Within the UK, urchin grazed biotopes (IR.MIR.KR.Lhyp.GzFt/Pk, IR.HIR.KFaR.LhypPar, IR.LIR.K.LhypSlat.Gz & IR.LIR.K.Slat.Gz) are generally localised to a few regions in North Scotland and Ireland (Smale *et al.*, 2013; Stenneck *et al.*, 2002; Norderhaug & Christie 2009; Connor *et al.*, 2004). IR.MIR.KR.Lhyp.GzFt/Pk, IR.HIR.KFaR.LhypPar, IR.LIR.K.LhypSlat.Gz & IR.LIR.K.Slat.Gz are characterized by a canopy-forming kelp. However, urchin grazing decreases the abundance and diversity of understory species. In the Isle of Man. Jones & Kain (1967) observed low *Echinus esculentus* grazing pressure can control the lower limit of *Laminaria hyperborea* and remove *Laminaria hyperborea* sporelings and juveniles. Urchin abundances in “urchin barrens” have been reported as high as 100 individuals/m<sup>2</sup> (Lang & Mann, 1978). Kain (1967) reported urchin abundances of 1-4/m<sup>2</sup> within experimental plots of the Isle of Man. Therefore, while “urchin barrens” are not presently an issue within the UK, relatively low urchin grazing has been found to control the depth distribution of *Laminaria hyperborea*, negatively impact on *Laminaria hyperborea* recruitment and reduce the understory community abundance and diversity.

Other factors that are likely to influence the recovery of *Laminaria hyperborea* biotopes is competitive interactions with Invasive Non-Indigenous Species (INIS), e.g. *Undaria pinnatifida* (Smale *et al.*, 2013; Brodie *et al.*, 2014; Heiser 2014), and/or the Lusitanian kelp *Laminaria ochroleuca* (Brodie *et al.*, 2014; Smale *et al.*, 2014). A predicted sea temperature rise in the North and Celtic seas of between 1.5-5°C over the next century (Philippart *et al.*, 2011) is likely to create northward range shifts in many macroalgal species, including *Laminaria hyperborea*. *Laminaria hyperborea* is a northern (Boreal) kelp species, thus increases in seawater temperature is likely to affect the resilience and recoverability of *Laminaria hyperborea* biotopes with southerly distributions in the UK (Smale *et al.*, 2013; Stenneck *et al.*, 2002). Evidence suggests that the Lusitanian kelp *Laminaria ochroleuca* (Smale *et al.*, 2014), and the INIS *Undaria pinnatifida* (Heiser *et al.*, 2014) are competing with *Laminaria hyperborea* along the UK south coast and may displace *Laminaria hyperborea* from some subtidal rocky reef habitats. The wider ecological consequences of *Laminaria hyperborea*' competition with *Laminaria ochroleuca* and *Undaria pinnatifida* are however as of yet unknown.

**Resilience assessment.** The evidence suggests that beds of mature *Laminaria hyperborea* can regenerate from disturbance within a period of 1-6 years, and the associated community within 7-10 years. However, other factors such as competitive interactions with *Laminaria ochroleuca* and *Undaria pinnatifida* may limit recovery of *Laminaria hyperborea* biotopes following disturbance. Also, urchin grazing pressure is shown to limit *Laminaria hyperborea* recruitment and reduce the diversity and abundance of the understory community and may limit habitat recovery following disturbance. The recovery of *Laminaria hyperborea* biotopes to disturbance from commercial harvesting in south Norway suggests that *Laminaria hyperborea* beds and the associated community could recover from a significant loss of canopy cover within 10 years, resilience has therefore been assessed as **Medium**.

Please note\* as in Northern Norway urchin grazing pressure could extend recovery/resilience of the *Laminaria hyperborea* biotopes >25 years, If intensive urchin grazing (as seen in Northern Norway) occurs in the UK resilience would be re-assessed as Very Low. However, because of the limited/localised incidence of urchin grazing within the UK, urchin grazing on large scales (as in Northern Norway) has not been included in this general resilience assessment. The introduction of Invasive Non-Indigenous Species (INIS) will also inhibit the recovery of *Laminaria hyperborea* biotopes for an indeterminate amount of time, in these cases, resilience would need to be re-assessed as Very Low. Another factor that is beyond the scope of this sensitivity assessment is the

presence of multiple concurrent synergistic or cumulative effects, which Smale *et al.* (2013) suggested could be a more damaging than individual pressures.

## Hydrological Pressures

	Resistance	Resilience	Sensitivity
Temperature increase (local)	Medium Q: High A: High C: High	Medium Q: High A: High C: High	Medium Q: High A: High C: High

Kain (1964) stated that *Laminaria hyperborea* sporophyte growth and reproduction could occur within a temperature range of 0 - 20°C. Upper and lower lethal temperatures have been estimated at between 1-2°C above or below the extremes of this range (Birkett *et al.*, 1988). Above 17°C gamete survival is reduced (Kain, 1964 & 1971) and gametogenesis is inhibited at 21°C (Dieck, 1992). It is therefore likely that *Laminaria hyperborea* recruitment will be impaired at a sustained temperature increase of above 17°C. Sporophytes however can tolerate slightly higher temperatures of 20°C. Temperature tolerances for *Laminaria hyperborea* are also seasonally variable and temperature changes are less tolerated in winter months than summer months (Birkett *et al.*, 1998).

Subtidal red algae are less tolerant of temperature extremes than intertidal red algae, surviving between -2°C and 18-23°C (Lüning 1990; Kain & Norton, 1990). Temperature increase may affect growth, recruitment or interfere with reproduction processes. For example, there is some evidence to suggest that blade growth in *Delesseria sanguinea* is delayed until ambient sea temperatures fall below 13°C. Blade growth is also likely to be intrinsically linked to gametangia development (Kain, 1987), and maintenance of sea temperatures above 13°C may affect recruitment success.

**Sensitivity assessment.** This biotope is distributed throughout the UK (Connor *et al.*, 2004). Northern to southern Sea Surface Temperature (SST) ranges from 8-16°C in summer and 6-13°C in winter (Beszczynska-Möller & Dye, 2013). Overall, a chronic change (2°C for a year) outside normal range for a year may reduce recruitment and growth, resulting in a minor loss in the population of kelp, especially in winter months or in southern examples of the biotope. However, an acute change (5°C for a month; e.g. from thermal effluent) may result in loss of abundance of kelp or extent of the bed, especially in winter. Therefore, resistance to the pressure is considered 'Medium', and resilience 'Medium'. The sensitivity of this biotope to increases in temperature has been assessed as 'Medium'.

	High	High	Not sensitive
Temperature decrease (local)	High Q: High A: High C: High	High Q: High A: High C: High	Not sensitive Q: High A: High C: High

Kain (1964) stated that *Laminaria hyperborea* sporophyte growth and reproduction could occur within a temperature range of 0 - 20°C. Upper and lower lethal temperatures have been estimated at between 1-2 °C above or below the extremes of this range (Birkett *et al.*, 1988). Subtidal red algae can survive at temperatures between -2 °C and 18-23 °C (Lüning, 1990; Kain & Norton, 1990).

*Laminaria hyperborea* is a boreal northern species with a geographic range from mid Portugal to Northern Norway (Birket *et al.*, 1998), and a mid range within southern Norway (60°-65° North)(Kain, 1971). The average seawater temperature for southern Norway in October is

12-13°C (Miller *et al.*, 2009), and average annual sea temperature, from 1970-2014, is 8°C (Beszczynska-Möller & Dye, 2013). The available information suggests that *Laminaria hyperborea* and biotope structure would not be affected by a change in sea temperature at the benchmark level.

**Sensitivity assessment.** Resistance to the pressure is considered 'High', and resilience 'High'. The sensitivity of this biotope to decreases in temperature has been assessed as 'Not Sensitive'.

**Salinity increase (local)**

<b>Low</b>	<b>Medium</b>	<b>Medium</b>
Q: Low A: NR C: NR	Q: High A: Medium C: High	Q: Low A: NR C: NR

Lüning (1990) suggest that "kelps" are stenohaline, their general tolerance to salinity as a phenotypic group covering 16 - 50 psu over a 24 hr period. Optimal growth probably occurs between 30-35 psu (MNCR category- 'Full' salinity) and growth rates are likely to be affected by periodic salinity stress. Birkett *et al.*, (1998) suggested that long-term increases in salinity may affect *Laminaria hyperborea* growth and may result in loss of affected kelp, and therefore loss of the biotope.

**Sensitivity assessment.** Resistance to the pressure is considered 'Low', and resilience 'Medium'. The sensitivity of this biotope to an increase in salinity has been assessed as 'Medium'.

**Salinity decrease (local)**

<b>Low</b>	<b>Medium</b>	<b>Medium</b>
Q: Medium A: Medium C: Medium	Q: High A: Medium C: High	Q: Medium A: Medium C: Medium

Lüning (1990) suggest that "kelps" are stenohaline, their general tolerance to salinity as a phenotypic group covering 16 - 50 psu over a 24 hr period. Optimal growth probably occurs between 30-35 psu (MNCR category-Full Salinity) and growth rates are likely to be affected by periodic salinity stress. Birkett *et al.*, (1998) suggest that long-term changes in salinity may result in loss of affected kelp and, therefore loss of this biotope.

Hopkin & Kain (1978) tested *Laminaria hyperborea* sporophyte growth at various low salinity treatments. The results showed that *Laminaria hyperborea* sporophytes could grow "normally" at 19 psu, growth was reduced at 16 psu and did not grow at 7 psu. A decrease in one MNCR salinity scale from 'Full' salinity (30-40psu) to 'Reduced' salinity (18-30 psu) would result in a decrease of *Laminaria hyperborea* sporophyte growth. *Laminaria hyperborea* may also be out-competed by low salinity tolerant species e.g. *Saccharina latissima* (Karsten, 2007), or the Invasive Non Indigenous Species *Undaria pinnatifida* (Burrows *et al.*, 2014).

If salinity was returned to 'Full' salinity (30-40 psu) *Laminaria hyperborea* could out-compete *Saccharina latissima* and re-establish community dominance in 2-4 years (Kain, 1975; Leinaas & Christie, 1996), however full habitat structure may take over 10 years to recover (Birkett *et al.*, 1998; Cristie *et al.*, 1998). The ability of *Laminaria hyperborea* to out-compete *Undaria pinnatifida* within the UK is however unknown (Heiser *et al.*, 2014), and as such interspecific interaction between *Laminaria hyperborea* and *Undaria pinnatifida* is not included within this sensitivity assessment.

**Sensitivity assessment.** Resistance to the pressure is considered 'Low', and resilience 'Medium'.

The sensitivity of this biotope to decreases in salinity has been assessed as 'Medium'.

### Water flow (tidal current) changes (local)

**High**

Q: High A: High C: High

**High**

Q: High A: High C: High

**Not sensitive**

Q: High A: High C: High

Kregting *et al.* (2013) measured *Laminaria hyperborea* blade growth and stipe elongation from an exposed and a sheltered site in Strangford Lough, Ireland, from March 2009-April 2010. Maximal significant wave height (Hm0) was 3.67 & 2m at the exposed and sheltered sites, and maximal water velocity (Velrms) was 0.6 & 0.3m/s at the exposed and sheltered sites respectively. Despite the differences in wave exposure and water velocity there was no significant difference in *Laminaria hyperborea* growth between the exposed and sheltered sites. Therefore water flow was found to have no significant effect on *Laminaria hyperborea* growth at the observed range of water velocities.

Biotope structure is however different between wave exposed and sheltered sites. Pederson *et al.* (2012) observed *Laminaria hyperborea* biomass, productivity and density increased with an increase in wave exposure. At low wave exposure *Laminaria hyperborea* canopy forming plants were smaller, had lower densities and had higher mortality rates than at exposed sites. At low wave exposure Pederson *et al.* (2012) suggested that high epiphytic loading on *Laminaria hyperborea* impaired light conditions, nutrient uptake, and increased the drag on the host *Laminaria hyperborea* during extreme storm events.

The morphology of the stipe and blade of kelps vary with water flow. In wave exposed areas, for example, *Laminaria hyperborea* develops a long and flexible stipe and this is probably a functional adaptation to strong water movement (Sjøtun, 1998). In addition, the lamina becomes narrower and thinner in strong currents (Sjøtun & Fredriksen, 1995). However, the stipe of *Laminaria hyperborea* is relatively stiff and can snap in strong currents. *Laminaria hyperborea* is usually absent from areas of high wave action or strong currents, although it is found in the Menai Strait, Wales, where tidal velocities can exceed 4 m/s (NBN, 2015) and in tidal rapids in Norway (J. Jones, pers. comm.) *Laminaria hyperborea* growth can persist in very strong tidal streams (>3 m/s).

Increase water flow rate may also remove or inhibit grazers including *Patella pellucida* and *Echinus esculentus* and remove epiphytic algae growth (Pederson *et al.*, 2012). The associated algal flora and suspension feeding faunal populations change significantly with different water flow regimes. Increased water flow rates may reduce the understory epiflora, to be replaced by an epifauna dominated community (e.g. sponges, anemones and polyclinid ascidians) as in the biotope IR.HIR.KFaR.LhypFa. The composition of the holdfast fauna may also change, e.g. energetic or sheltered water movements favour different species of amphipods (Moore, 1985).

IR.HIR.KFaR.LhypR, IR.HIR.KFaR.LhypFa, IR.MIR.KR.Lhyp, and their associated sub-biotopes are found within strong (1.5-3 m/s)-moderate (0.5-1.5 m/s) tidal streams. A change in peak mean spring bed flow velocity which does not result in a change in tidal streams above or below 0.5-3 m/s is not likely to affect the dominance of *Laminaria hyperborea* within the community, but may cause changes in the understory community. The prominent understory filter feeding community within IR.HIR.KFaR.LhypFa is reliant on high water movement. A decrease in tidal streams may result in a decline of filter feeding fauna and an increase in red seaweeds within the understory community or vice versa with an increase in tidal streams A decrease in tidal flow within this range may also decrease urchin dislodgment and increase urchin grazing. An increase in urchin grazing may cause a decline in the understory community abundance and diversity (as in IR.MIR.KR.Lhyp.GzFt/Pk and IR.MIR.KR.LhypPar).

**Sensitivity assessment.** A change in peak mean spring bed flow velocity of between 0.1m/s to 0.2m/s for more than 1 year is not likely to affect the dominance of *Laminaria hyperborea*, however subtle differences in tidal regime may influence the understory community. Resistance to the pressure is considered 'High', and resilience 'High'. Hence, the sensitivity of this biotope to changes in peak mean spring bed velocity has been assessed as 'Not Sensitive'.

However, if peak mean spring bed flow velocity changes but remains within 0.5-3 m/s *Laminaria hyperborea* is likely to remain the dominant habitat but the understory community may be affected; directly by a change in water velocity or through increased grazing pressure.

### Emergence regime changes

**Low**

Q: Low A: NR C: NR

**Medium**

Q: High A: Low C: High

**Medium**

Q: Low A: NR C: NR

The upper limit of the *Laminaria hyperborea* bed is determined by wave action and water flow, desiccation, and competition from the more emergence resistant *Laminaria digitata*. *Laminaria hyperborea* exposed at extreme low water are very intolerant of desiccation, the most noticeable effect being bleaching of the frond and subsequent death of the meristem and loss of the plant. An increase in wave exposure (see below- water flow), as a result of increased emergence, has been found to exclude *Laminaria hyperborea* from shallow waters due to dislodgement of the sporophyte or snapping of the stipe (Birket *et al.*, 1998). Hence, an increase in emergence is likely to lead to mortality of exposed *Laminaria hyperborea* and the associated habitat.

An increase in water depth/decreased emergence (at the benchmark level) may increase the upper depth restriction of *Laminaria hyperborea* forest biotope variants. However, limited light availability at depth will decrease the lower extent of *Laminaria hyperborea*, and may therefore result in a shift from forest to park biotope variants at depth. Further increases in depth will cause a community shift to that characterized by circalittoral faunal species, however this is beyond the scope of the benchmark.

**Sensitivity assessment.** Resistance to the pressure is considered 'Low', and resilience 'Medium'. The sensitivity of this biotope to changes in tidal emergence has been assessed as 'Medium'.

### Wave exposure changes (local)

**High**

Q: High A: High C: High

**High**

Q: High A: High C: High

**Not sensitive**

Q: High A: High C: High

Kregting *et al.*, (2013) measured *Laminaria hyperborea* blade growth and stipe elongation from an exposed and a sheltered site in Strangford Lough, Ireland from March 2009-April 2010. Wave exposure was found to be between 1.1. to 1.6 times greater between the exposed and sheltered sites. Maximal significant wave height (Hm0) was 3.67 & 2m at the exposed and sheltered sites. Maximal water velocity (Velrms) was 0.6 & 0.3m/s at the exposed and sheltered sites. Despite the differences in wave exposure and water velocity there was no significant difference in *Laminaria hyperborea* growth between the exposed and sheltered site.

Biotope structure is however different between wave exposed and sheltered sites. Pederson *et al.*, (2012) observed *Laminaria hyperborea* biomass, productivity and density increased with an increase in wave exposure. At low wave exposure *Laminaria hyperborea* canopy forming plants were smaller, had lower densities and had higher mortality rates than at exposed sites. At low

wave exposure high epiphytic loading on *Laminaria hyperborea* was theorised to impair light conditions, nutrient uptake, and increase the drag of the host *Laminaria hyperborea* during extreme storm events.

The morphology of the stipe and blade of kelps vary with water flow. In wave exposed areas, for example, *Laminaria hyperborea* develops a long and flexible stipe and this is probably a functional adaptation to strong water movement (Sjøtun, 1998). In addition, the lamina becomes narrower and thinner in strong currents (Sjøtun & Fredriksen, 1995). However, the stipe of *Laminaria hyperborea* is relatively stiff and can snap in strong currents. *Laminaria hyperborea* is usually absent from areas of extreme wave action and can be replaced by *Alaria esculenta*. In extreme wave exposures *Alaria esculenta* can dominate the shallow sub-littoral to a depth of 15m (Birket *et al.*, 1998).

Increase water flow rate may also remove or inhibit grazers including *Patella pellucida* and *Echinus esculentus* and remove epiphytic algae growth (Pederson *et al.*, 2012). The associated algal flora and suspension feeding faunal populations change significantly with different water flow regimes. Increased water flow rates may reduce the understory epiflora, to be replaced by an epifauna dominated community (e.g. sponges, anemones and polyclinid ascidians) as in the biotope IR.HIR.KFaR.LhypFa. The composition of the holdfast fauna may also change, e.g. energetic or sheltered water movements favour different species of amphipods (Moore, 1985).

IR.HIR.KFaR.LhypR, IR.HIR.KFaR.LhypFa, IR.MIR.KR.Lhyp, and their associated sub-biotopes are found between extremely exposed to moderate wave exposure. Changes in local wave height above or below that experienced in extremely exposed to moderately exposed sites will affect the dominance of *Laminaria hyperborea*. Smaller changes in local wave height have the potential to cause changes to the understory community. The prominent understory filter feeding community within IR.HIR.KFaR.LhypFa is reliant on wave surge currents. A decrease in wave surge may result in a decline of filter feeding fauna and an increase in red seaweeds within the understory community or vice versa. A decrease in local wave height may also decrease the chance of urchins being dislodged (removed) from biotopes found at sites with traditionally high wave exposure and may therefore increase urchin grazing. An increase in urchin grazing may cause a decline in the understory community abundance and diversity (as in IR.MIR.KR.Lhyp.GzFt/Pk and IR.MIR.KR.LhypPar).

**Sensitivity assessment.** A change in nearshore significant wave height >3% but <5% is however unlikely to have a significant effect. Resistance to the pressure is considered 'High', and resilience 'High'. Hence, the sensitivity of this biotope to changes in local wave height has been assessed as 'Not Sensitive'.

## Chemical Pressures

	Resistance	Resilience	Sensitivity
Transition elements & organo-metal contamination	Not Assessed (NA) Q: NR A: NR C: NR	Not assessed (NA) Q: NR A: NR C: NR	Not assessed (NA) Q: NR A: NR C: NR

This pressure is **Not assessed** but evidence is presented where available.

Bryan (1984) suggested that the general order for heavy metal toxicity in seaweeds is: Organic Hg

> inorganic Hg > Cu > Ag > Zn > Cd > Pb. Cole *et al.*, (1999) reported that Hg was very toxic to macrophytes. Similarly, Hopkin & Kain (1978) demonstrated sub-lethal effects of heavy metals on *Laminaria hyperborea* gametophytes and sporophytes, including reduced growth and respiration. Sheppard *et al.*, (1980) noted that increasing levels of heavy metal contamination along the west coast of Britain reduced species number and richness in holdfast fauna, except for suspension feeders which became increasingly dominant. Gastropods may be relatively tolerant of heavy metal pollution (Bryan, 1984). *Echinus esculentus* recruitment is likely to be impaired by heavy metal contamination due to the intolerance of its larvae. *Echinus esculentus* are long-lived and poor recruitment may not reduce grazing pressure in the short-term. Although macroalgae species may not be killed, except by high levels of contamination, reduced growth rates may impair the ability of the biotope to recover from other environmental disturbances.

### Hydrocarbon & PAH contamination

Not Assessed (NA)

Q: NR A: NR C: NR

Not assessed (NA)

Q: NR A: NR C: NR

Not assessed (NA)

Q: NR A: NR C: NR

This pressure is **Not assessed** but evidence is presented where available.

*Laminaria hyperborea* fronds, being almost exclusively sub tidal, would not come into contact with freshly released oil, but only to sinking emulsified oil and oil adsorbed onto particles (Birket *et al.*, 1998). The mucilaginous slime layer coating of laminarians may protect them from smothering by oil. Hydrocarbons in solution reduce photosynthesis and may be algicidal. However, Holt *et al.*, (1995) reported that oil spills in the USA and from the 'Torrey Canyon' had little effect on kelp forests. Similarly, surveys of subtidal communities at a number sites between 1-22.5m below chart datum, including *Laminaria hyperborea* communities, showed no noticeable impacts of the Sea Empress oil spill and clean up (Rostron & Bunker, 1997). An assessment of holdfast fauna in *Laminaria* showed that although species richness and diversity decreased with increasing proximity to the Sea Empress oil spill, overall the holdfasts contained a reasonably rich and diverse fauna, even though oil was present in most samples (Sommerfield & Warwick, 1999). Laboratory studies of the effects of oil and dispersants on several red algae species, including *Delesseria sanguinea* (Grandy 1984; cited in Holt *et al.*, 1995) concluded that they were all sensitive to oil/dispersant mixtures, with little differences between adults, sporelings, diploid or haploid life stages. Holt *et al.*, (1995) concluded that *Delesseria sanguinea* is probably generally sensitive of chemical contamination. Overall the red algae are likely to be highly intolerant to hydrocarbon contamination. Loss of red algae is likely to reduce the species richness and diversity of the biotope and the understory may become dominated by encrusting corallines; however, red algae are likely to recover relatively quickly.

### Synthetic compound contamination

Not Assessed (NA)

Q: NR A: NR C: NR

Not assessed (NA)

Q: NR A: NR C: NR

Not assessed (NA)

Q: NR A: NR C: NR

This pressure is **Not assessed** but evidence is presented where available.

O'Brian & Dixon (1976) suggested that red algae were the most sensitive group of macrophytes to oil and dispersant contamination (see Smith, 1968). Although *Laminaria hyperborea* sporelings and gametophytes are intolerant of atrazine (and probably other herbicides) overall they may be relatively tolerant of synthetic chemicals (Holt *et al.*, 1995). *Laminaria hyperborea* survived within >55m from the acidified halogenated effluent discharge polluting Amlwch Bay, Anglesey, albeit at low density. These specimens were greater than 5 years of age, suggesting that spores and/or early stages were more intolerant (Hoare & Hiscock, 1974). *Patella pellucida* was excluded from Amlwch

Bay by the pollution and the species richness of the holdfast fauna decreased with proximity to the effluent discharge; amphipods were particularly intolerant although polychaetes were the least affected (Hoare & Hiscock, 1974). The richness of epifauna/flora decreased near the source of the effluent and epiphytes were absent from *Laminaria hyperborea* stipes within Amlwch Bay. The red alga *Phyllophora membranifolia* was also tolerant of the effluent in Amlwch Bay. Smith (1968) also noted that epiphytic and benthic red algae were intolerant of dispersant or oil contamination due to the Torrey Canyon oil spill; only the epiphytes *Cryptopleura ramosa* and *Spermothamnion repens* and some tufts of *Jania rubens* survived together with *Osmundea pinnatifida*, *Gigartina pistillata* and *Phyllophora crispa* from the sublittoral fringe. *Delesseria sanguinea* was probably to most intolerant since it was damaged at depths of 6m (Smith, 1968). Holt *et al.*, (1995) suggested that *Delesseria sanguinea* is probably generally sensitive of chemical contamination. Although *Laminaria hyperborea* may be relatively insensitive to synthetic chemical pollution, evidence suggests that grazing gastropods, amphipods and red algae are sensitive. Loss of red algae is likely to reduce the species richness and diversity of the biotope and the understorey may become dominated by encrusting corallines; however, red algae are likely to recover relatively quickly.

#### Radionuclide contamination

Not relevant (NR)

Q: NR A: NR C: NR

Not relevant (NR)

Q: NR A: NR C: NR

No evidence (NEv)

Q: NR A: NR C: NR

No evidence

#### Introduction of other substances

Not Assessed (NA)

Q: NR A: NR C: NR

Not assessed (NA)

Q: NR A: NR C: NR

Not assessed (NA)

Q: NR A: NR C: NR

This pressure is **Not assessed**.

#### De-oxygenation

High

Q: High A: Medium C: High

High

Q: High A: Medium C: High

Not sensitive

Q: High A: Medium C: High

Reduced oxygen concentrations have been shown to inhibiting both photosynthesis and respiration in macroalgae (Kinne, 1977). Despite this, macroalgae are thought to buffer the environmental conditions of low oxygen, thereby acting as a refuge for organisms in oxygen depleted regions especially if the oxygen depletion is short-term (Frieder *et al.*, 2012). A rapid recovery from a state of low oxygen is expected if the environmental conditions are transient. If levels do drop below 4 mg/l negative effects on these organisms can be expected with adverse effects occurring below 2mg/l (Cole *et al.*, 1999).

**Sensitivity Assessment.** Reduced oxygen levels are likely to inhibit photosynthesis and respiration but not cause a loss of the macroalgae population directly. However, small invertebrate epifauna may be lost, causing a reduction in species richness. Therefore a resistance of 'High' is recorded. Resilience is likely to be 'High', and the biotopes is probably 'Not sensitive' at the benchmark level.

#### Nutrient enrichment

Not relevant (NR)

Q: NR A: NR C: NR

Not relevant (NR)

Q: NR A: NR C: NR

Not sensitive

Q: NR A: NR C: NR

Holt *et al.* (1995) suggest that *Laminaria hyperborea* may be tolerant of nutrient enrichment since healthy populations are found at ends of sublittoral untreated sewage outfalls in the Isle of Man. Increased nutrient levels e.g. from sewage outfalls, has been associated with increases in

abundance, primary biomass and *Laminaria hyperborea* stipe production but with concomitant decreases in species numbers and diversity (Fletcher, 1996).

Increased nutrients may result in phytoplankton blooms that increase turbidity (see above). Increased nutrients may favour sea urchins, e.g. *Echinus esculentus*, due their ability to absorb dissolved organics, and result in increased grazing pressure leading to loss of understorey epiflora/fauna, decreased kelp recruitment and possibly 'urchin barrens'. Therefore, although nutrients may not affect kelps directly, indirect effects such as turbidity, siltation and competition may significantly affect the structure of the biotope.

However this biotope is considered to be '**Not sensitive**' at the pressure benchmark, that assumes compliance with good status as defined by the WFD.

<b>Organic enrichment</b>	<b>Medium</b>	<b>High</b>	<b>Low</b>
	Q: Medium A: Medium C: Medium	Q: High A: Medium C: High	Q: Medium A: Medium C: Medium

Holt *et al.* (1995) suggest that *Laminaria hyperborea* may be tolerant of organic enrichment since healthy populations are found at ends of sub littoral untreated sewage outfalls in the Isle of Man. Increased nutrient levels e.g. from sewage outfalls, has been associated with increases in abundance, primary biomass and *Laminaria hyperborea* stipe production but with concomitant decreases in species numbers and diversity (Fletcher, 1996). Increase in ephemeral and opportunistic algae are associated with reduced numbers of perennial macrophytes (Fletcher, 1996). Increased nutrients may also result in phytoplankton blooms that increase turbidity. Therefore, although nutrients may not affect kelps directly, indirect effects such as turbidity may significantly affect the structure of *Laminaria hyperborea* biotopes.

**Sensitivity assessment.** Resistance to the pressure is considered '**Medium**', and resilience '**High**'. The sensitivity of this biotope to organic enrichment is assessed as '**Low**'.

## **A Physical Pressures**

	<b>Resistance</b>	<b>Resilience</b>	<b>Sensitivity</b>
<b>Physical loss (to land or freshwater habitat)</b>	<b>None</b>	<b>Very Low</b>	<b>High</b>
	Q: High A: High C: High	Q: High A: High C: High	Q: High A: High C: High

All marine habitats and benthic species are considered to have a resistance of 'None' to this pressure and to be unable to recover from a permanent loss of habitat (resilience is '**Very Low**'). Sensitivity within the direct spatial footprint of this pressure is therefore '**High**'. Although no specific evidence is described confidence in this assessment is '**High**', due to the incontrovertible nature of this pressure.

<b>Physical change (to another seabed type)</b>	<b>None</b>	<b>Very Low</b>	<b>High</b>
	Q: High A: High C: High	Q: High A: High C: High	Q: High A: High C: High

If rock substrata were replaced with sedimentary substrata this would represent a fundamental change in habitat type, which *Laminaria hyperborea* would not be able to tolerate (Birket *et al.*, 1998). The biotope would be lost.

**Sensitivity assessment.** Resistance to the pressure is considered “None”, and resilience “Very Low” or ‘None’. The sensitivity of this biotope to change from sedimentary or soft rock substrata to hard rock or artificial substrata or vice-versa is assessed as “High”.

<b>Physical change (to another sediment type)</b>	Not relevant (NR) Q: NR A: NR C: NR	Not relevant (NR) Q: NR A: NR C: NR	Not relevant (NR) Q: NR A: NR C: NR
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Not relevant

<b>Habitat structure changes - removal of substratum (extraction)</b>	Not relevant (NR) Q: NR A: NR C: NR	Not relevant (NR) Q: NR A: NR C: NR	Not relevant (NR) Q: NR A: NR C: NR
-----------------------------------------------------------------------	----------------------------------------	----------------------------------------	----------------------------------------

Not relevant to rock substrata.

<b>Abrasion/disturbance of the surface of the substratum or seabed</b>	Low Q: High A: High C: High	Medium Q: High A: High C: High	Medium Q: High A: High C: High
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Christie *et al.*, (1998) observed *Laminaria hyperborea* habitat regeneration following commercial *Laminaria hyperborea* trawling in south Norway. Within the study area, trawling removed all large canopy-forming adult *Laminaria hyperborea*, however sub-canopy recruits were largely unaffected. In 2-6 years of harvesting a new canopy had formed 1m off the seabed. The associated holdfast communities recovered in 6 years, however the epiphytic stipe community did not fully recover within the same time period. Christie *et al.*, (1998) suggested that kelp habitats were relatively resistant to direct disturbance/removal of *Laminaria hyperborea* canopy.

Recurrent disturbance occurring at a smaller time scale than the recovery period of 2-6 years (stated above) could extend recovery time. Kain (1975) cleared sublittoral blocks of *Laminaria hyperborea* at different times of the year for several years. The first colonizers and succession community differed between blocks and at what time of year the blocks were cleared however within 2 years of clearance the blocks were dominated by *Laminaria hyperborea*. Lienaaas & Christie (1996) also observed *Laminaria hyperborea* re-colonization of “urchin barrens”, following removal of urchins. The substratum was initially colonized by filamentous macroalgae and *Saccharina latissima* however after 2-4 years *Laminaria hyperborea* dominated the community.

**Sensitivity assessment.** Resistance to the pressure is considered ‘Low’, and resilience ‘Medium’. The sensitivity of this biotope to damage to seabed surface features is assessed as ‘Medium’.

<b>Penetration or disturbance of the substratum subsurface</b>	Not relevant (NR) Q: NR A: NR C: NR	Not relevant (NR) Q: NR A: NR C: NR	Not relevant (NR) Q: NR A: NR C: NR
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Not relevant, please refer to pressure “Abrasion/disturbance of the substrata on the surface of the seabed”.

**Changes in suspended solids (water clarity)****Low**

Q: High A: High C: High

**Medium**

Q: High A: High C: High

**Medium**

Q: High A: High C: High

Suspended Particle Matter (SPM) concentration has a linear relationship with sub-surface light attenuation (Kd) (Devlin *et al.*, 2008). An increase in SPM results in a decrease in sub-surface light attenuation. Light availability and water turbidity are principal factors in determining kelp depth range (Birkett *et al.*, 1998). Light penetration influences the maximum depth at which kelp species can grow and it has been reported that laminarians grow down to depths at which the light levels are reduced to 1 percent of incident light at the surface. Maximal depth distribution of laminarians, therefore, varies from 100 m in the Mediterranean to only 6-7 m in the silt-laden German Bight. In Atlantic European waters, the depth limit is typically 35 m. In very turbid waters the depth at which *Laminaria hyperborea* is found may be reduced, or in some cases excluded completely (e.g. Severn Estuary), because of the alteration in light attenuation by suspended sediment (Birkett *et al.* 1998b; Lüning, 1990).

*Laminaria* spp. show a decrease of 50% photosynthetic activity when turbidity increases by 0.1/m (light attenuation coefficient =0.1-0.2/m; Staehr & Wernberg, 2009). An increase in water turbidity will likely affect the photosynthetic ability of *Laminaria hyperborea* and *Laminaria ochroleuca* and decrease *Laminaria hyperborea* abundance and density (see sub-biotope-IR.MIR.KR.Lhyp.Pk). Kain (1964) suggested that early *Laminaria hyperborea* gametophyte development could occur in the absence of light. Furthermore, observations from south Norway found that a pool of *Laminaria hyperborea* recruits could persist growing beneath *Laminaria hyperborea* canopies for several years, indicating that sporophyte growth can occur in light-limited environments (Christe *et al.*, 1998). However in habitats exposed to high levels of suspended silts *Laminaria hyperborea* is out-competed by *Saccharina latissima*, a silt tolerant species, and thus, a decrease in water clarity is likely to decrease the abundance of *Laminaria hyperborea* in the affected area (Norton, 1978).

**Sensitivity Assessment.** Changes in water clarity are likely to affect photosynthetic rates and enable *Saccharina latissima* to compete more successfully with *Laminaria hyperborea*. A decrease in turbidity is likely to support enhanced growth (and possible habitat expansion) and is therefore not considered in this assessment. An increase in water clarity from clear to intermediate (10-100 mg/l) represents a change in light attenuation of ca 0.67-6.7 Kd/m, and is likely to result in a greater than 50% reduction in photosynthesis of *Laminaria* spp. Therefore, the dominant kelp species will probably suffer a significant decline and resistance to this pressure is assessed as 'Low'. Resilience to this pressure is probably 'Medium' at the benchmark. Hence, this biotope is assessed as having a sensitivity of 'Medium' to this pressure.

**Smothering and siltation rate changes (light)****High**

Q: Low A: NR C: NR

**High**

Q: Low A: NR C: NR

**Not sensitive**

Q: Low A: NR C: NR

Smothering by sediment e.g. 5 cm material during a discrete event, is unlikely to damage *Laminaria hyperborea* sporophytes but is likely to affect gametophyte survival as well as holdfast fauna, and interfere with zoospore settlement. Given the microscopic size of the gametophyte, 5 cm of sediment could be expected to significantly inhibit growth. However, laboratory studies showed that gametophytes can survive in darkness for between 6 - 16 months at 8 °C and would probably survive smothering by a discrete event. Once returned to normal conditions the gametophytes resumed growth or maturation within one month (Dieck, 1993). Intolerance to this factor is likely to be higher during the peak periods of sporulation and/or spore settlement.

If inundation is long lasting then the understory epifauna/flora may be adversely affected, e.g. suspension or filter feeding fauna and/or algal species. This biotope occurs in high wave exposures and therefore deposited sediments are unlikely to remain for more than a few tidal cycles, except in the deepest of rock-pools. Therefore the effects of depositing 5 cm of fine sediment in a discrete event are likely to be transient.

**Sensitivity assessment.** Resistance to the pressure is considered '**High**', and resilience '**High**'. The sensitivity of this biotope to light deposition of up to 5 cm of fine material added to the seabed in a single discrete event is assessed as '**Note Sensitive**'.

### Smothering and siltation rate changes (heavy)

**Medium**

Q: **Low** A: **NR** C: **NR**

**High**

Q: **Low** A: **NR** C: **NR**

**Low**

Q: **Low** A: **NR** C: **NR**

Smothering by sediment e.g. 30 cm material during a discrete event, is unlikely to damage *Laminaria hyperborea* plants but is likely to affect gametophyte survival, holdfast communities, epiphytic community at the base of the stipe, and interfere with zoospore settlement. Given the microscopic size of the gametophyte, 30 cm of sediment could be expected to significantly inhibit growth. However, laboratory studies showed that gametophytes can survive in darkness for between 6 - 16 months at 8 °C and would probably survive smothering within a discrete event. Once returned to normal conditions the gametophytes resumed growth or maturation within one month (Dieck, 1993). Intolerance to this factor is likely to be higher during the peak periods of sporulation and/or spore settlement.

If clearance of deposited sediment occurs rapidly then understory communities are expected to recover quickly. If inundation is long lasting then the understory epifauna/flora may be adversely affected, e.g. suspension or filter feeding fauna and/or algal species. While this biotope occurs in high to moderate energy habitats (due to water flow or wave action) deposition of 30cm of sediment represents a large volume of material that would likely remain for a number of tidal cycles and is expected to damage understory flora/fauna as well as juvenile *Laminaria hyperborea*.

**Sensitivity assessment.** Resistance to the pressure is considered '**Medium**', and resilience '**High**'. The sensitivity of this biotope to heavy deposition of up to 30cm of fine material added to the seabed in a single discrete event is assessed as '**Low**'.

### Litter

**Not Assessed (NA)**

Q: **NR** A: **NR** C: **NR**

**Not assessed (NA)**

Q: **NR** A: **NR** C: **NR**

**Not assessed (NA)**

Q: **NR** A: **NR** C: **NR**

Not assessed.

### Electromagnetic changes

**Not relevant (NR)**

Q: **NR** A: **NR** C: **NR**

**Not relevant (NR)**

Q: **NR** A: **NR** C: **NR**

**No evidence (NEv)**

Q: **NR** A: **NR** C: **NR**

No evidence

### Underwater noise changes

**Not relevant (NR)**

Q: **NR** A: **NR** C: **NR**

**Not relevant (NR)**

Q: **NR** A: **NR** C: **NR**

**Not relevant (NR)**

Q: **NR** A: **NR** C: **NR**

Not relevant

**Introduction of light or shading****Low**

Q: Low A: NR C: NR

**Medium**

Q: Low A: NR C: NR

**Medium**

Q: Low A: Low C: Low

Shading of the biotope (e.g. by construction of a pontoon, pier etc) could adversely affect the biotope in areas where the water clarity is also low, and tip the balance to shade tolerant species, resulting in the loss of the biotope directly within the shaded area, or a reduction in laminarian abundance from forest to park type biotopes.

**Sensitivity assessment.** Resistance is probably 'Low', with a 'Medium' resilience and a sensitivity of 'Medium', albeit with 'low' confidence due to the lack of direct evidence.

**Barrier to species movement****Not relevant (NR)**

Q: NR A: NR C: NR

**Not relevant (NR)**

Q: NR A: NR C: NR

**Not relevant (NR)**

Q: NR A: NR C: NR

**Not relevant.** This pressure is considered applicable to mobile species, e.g. fish and marine mammals rather than seabed habitats. Physical and hydrographic barriers may limit the dispersal of spores. But spore dispersal is not considered under the pressure definition and benchmark.

**Death or injury by collision****Not relevant (NR)**

Q: NR A: NR C: NR

**Not relevant (NR)**

Q: NR A: NR C: NR

**Not relevant (NR)**

Q: NR A: NR C: NR

**Not relevant.** Collision from grounding vessels is addressed under abrasion above.

**Visual disturbance****Not relevant (NR)**

Q: NR A: NR C: NR

**Not relevant (NR)**

Q: NR A: NR C: NR

**Not relevant (NR)**

Q: NR A: NR C: NR

Not relevant

**Biological Pressures****Resistance****Resilience****Sensitivity****Genetic modification & translocation of indigenous species****Not relevant (NR)**

Q: NR A: NR C: NR

**Not relevant (NR)**

Q: NR A: NR C: NR

**No evidence (NEV)**

Q: NR A: NR C: NR

At the time of writing **no evidence** regarding the genetic modification or effects of translocation of native kelp populations was found.

**Introduction or spread of invasive non-indigenous species****Low**

Q: High A: High C: High

**Very Low**

Q: High A: High C: High

**High**

Q: High A: High C: High

*Undaria pinnatifida* has received a large amount of research attention as a major Invasive Non Indigenous Species (INIS) which could out-compete native UK kelp habitats (see Farrell & Fletcher, 2006; Thompson & Schiel, 2012, Brodie *et al.*, 2014; Hieser *et al.*, 2014). *Undaria pinnatifida* was first recorded in Plymouth Sound, UK in 2003 (NBN, 2015) subsequent surveys in 2011 have

reported that *Undaria pinnatifida* is widespread throughout Plymouth Sound, colonizing rocky reef habitats. Where *Undaria pinnatifida* is present there was a significant decrease in the abundance of other *Laminaria* species, including *Laminaria hyperborea* (Heiser *et al.*, 2014).

In New Zealand, Thompson & Schiel (2012) observed that native fucoids could out-compete *U. pinnatifida* and re-dominate the substratum. However, Thompson & Schiel (2012) suggested the fucoid recovery was partially due to an annual *Undaria pinnatifida* die back, which as noted by Heiser *et al.*, (2014) did not occur in Plymouth Sound, UK. It is unknown whether *Undaria pinnatifida* will out-compete native macro-algae in the UK. However, from 2003-2011 *Undaria pinnatifida* had spread throughout Plymouth Sound, UK, becoming a visually dominant species at some locations within summer months (Hieser *et al.*, 2014). While *Undaria pinnatifida* may replace *Laminaria hyperborea* in some locations within the UK, at the time of writing there is limited evidence available to assess what ecological impacts this invasion may have on *Laminaria hyperborea* associated communities e.g. red seaweeds.

*Undaria pinnatifida* was successfully eradicated on a sunken ship in Clatham Islands, New Zealand, by applying a heat treatment of 70 °C (see Wotton *et al.*, 2004) however numerous other eradication attempts have failed, and as noted by Farrell & Fletcher (2006) once established *Undaria pinadifida* resists most attempts of long-term removal. The biotope is unlikely to fully recover until *Undaria pinnatifida* is fully removed from the habitat, which as stated above is unlikely to occur.

**Sensitivity assessment.** Resistance to the pressure is considered 'Low', and resilience 'Very Low'. The sensitivity of this biotope to introduction of INIS is assessed as 'High'.

#### Introduction of microbial pathogens

Medium

Q: Low A: NR C: NR

High

Q: High A: Low C: High

Low

Q: Low A: NR C: NR

Galls on the blade of *Laminaria hyperborea* and spot disease are associated with the endophyte *Streblonema* sp. although the causal agent is unknown (bacteria, virus or endophyte). Resultant damage to the blade and stipe may increase losses in storms. The endophyte inhibits spore production and therefore recruitment and recoverability (Lein *et al.*, 1991).

**Sensitivity assessment.** Resistance to the pressure is considered 'Medium', and resilience 'High'. The sensitivity of this biotope to introduction of microbial pathogens is assessed as 'Low'.

#### Removal of target species

None

Q: High A: High C: High

Medium

Q: High A: High C: High

Medium

Q: High A: High C: High

Christie *et al.* (1998) observed *Laminaria hyperborea* habitat regeneration following commercial *Laminaria hyperborea* trawling in south Norway. Within the study area trawling removed all large canopy-forming adult *Laminaria hyperborea*, however sub-canopy recruits were unaffected. Within 2-3 years of harvesting a new canopy had formed 1 m off the seabed. The associated holdfast communities recovered in 6 years however the epiphytic stipe community did not fully recover within the same time period. Christie *et al.*, (1998) suggested that kelp habitats were relatively resistant to direct disturbance of *Laminaria hyperborea* canopy.

Recurrent disturbance occurring at a smaller time scale than the recovery period of 2-6 years (stated above) could extend recovery time. Kain (1975) cleared sublittoral blocks of *Laminaria*

*hyperborea* at different times of the year for several years. The first colonizers and succession community differed between blocks and at what time of year the blocks were cleared however within 2 years of clearance the blocks were dominated by *Laminaria hyperborea*. Lienaas & Christie (1996) also observed *Laminaria hyperborea* re-colonization of “urchin barrens”, following removal of urchins. The substratum was initially colonized by filamentous macroalgae and *Saccharina latissima* however after 2-4 years *Laminaria hyperborea* dominated the community.

Following disturbance or in areas where recurrent rapid disturbance occurs *Laminaria hyperborea* recruitment could also be affected by interspecific competitive interactions with Invasive Non Indigenous Species or ephemeral algal species (Brodie *et al.*, 2013; Smale *et al.*, 2013), however evidence for this is limited and thus not included within this assessment.

**Sensitivity assessment.** Resistance to the pressure is considered ‘None’, and resilience ‘Medium’. The sensitivity of this biotope to damage to seabed surface features is assessed as ‘Medium’.

#### Removal of non-target species

**Low**

Q: High A: High C: High

**Medium**

Q: High A: High C: High

**Medium**

Q: High A: High C: High

Incidental/accidental removal of *Laminaria hyperborea* from extraction of other marine resources, e.g. fisheries or aggregates, is likely to cause similar effects to that of direct harvesting of *Laminaria hyperborea*; hence the same evidence has been used for both pressure assessments.

Christie *et al.* (1998) observed *Laminaria hyperborea* habitat regeneration following commercial *Laminaria hyperborea* trawling in south Norway. Within the study area trawling removed all large canopy-forming adult *Laminaria hyperborea*, however sub-canopy recruits were unaffected. Within 2-6 years of harvesting a new canopy had formed 1m off the seabed. The associated holdfast communities recovered in 6 years however the epiphytic stipe community did not fully recover within the same time period. Christie *et al.*, (1998) suggested that kelp habitats were relatively resistant to direct disturbance of *Laminaria hyperborea* canopy.

Recurrent disturbance occurring at a smaller time scale than the recovery period of 2-6 years (stated above) could extend recovery time. Kain (1975) cleared sublittoral blocks of *Laminaria hyperborea* at different times of the year for several years. The first colonizers and succession community differed between blocks and at what time of year the blocks were cleared however within 2 years of clearance the blocks were dominated by *Laminaria hyperborea*. Lienaas & Christie (1996) also observed *Laminaria hyperborea* re-colonization of “urchin barrens”, following removal of urchins. The substratum was initially colonized by filamentous macroalgae and *Saccharina latissima* however after 2-4 years *Laminaria hyperborea* dominated the community.

Following disturbance or in areas where recurrent rapid disturbance occurs *Laminaria hyperborea* recruitment could also be affected by interspecific competitive interactions with Invasive Non Indigenous Species or ephemeral algal species (Brodie *et al.*, 2013; Smale *et al.*, 2013), however evidence for this is limited and thus not included within this assessment.

**Sensitivity assessment.** Resistance to the pressure is considered ‘Low’, and resilience ‘Medium’. The sensitivity of this biotope to damage to seabed surface features is assessed as ‘Medium’.

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