# ICES Journal of <br> Marine Science 

## Original Article

# Estimating biomass, fishing mortality, and "total allowable discards" for surveyed non-target fish 

Samuel Shephard ${ }^{1 *}$, David G. Reid ${ }^{2}$, Hans D. Gerritsen ${ }^{2}$, and Keith D. Farnsworth ${ }^{1}$<br>${ }^{1}$ School of Biological Sciences, Queen's University Belfast, 97 Lisburn Road, Belfast BT9 7BL, UK<br>${ }^{2}$ Marine Institute, Rinville, Oranmore, Co. Galway, Ireland<br>*Corresponding author: e-mail: s.shephard@qub.ac.uk<br>Shephard, S., Reid, D. G., Gerritsen, H. D., and Farnsworth, K. D. Estimating biomass, fishing mortality, and "total allowable discards" for surveyed non-target fish. - ICES Journal of Marine Science, 72: 458-466.

Received 1 April 2014; revised 29 July 2014; accepted 31 July 2014; advance access publication 27 August 2014.
Demersal fisheries targeting a few high-value species often catch and discard other "non-target" species. It is difficult to quantify the impact of this incidental mortality when population biomass of a non-target species is unknown. We calculate biomass for 14 demersal fish species in ICES Area VIIg (Celtic Sea) by applying species- and length-based catchability corrections to catch records from the Irish Groundfish Survey (IGFS). We then combine these biomass estimates with records of commercial discards (and landings for marketable non-target species) to calculate annual harvesting rates $(H R)$ for each study species. Uncertainty is incorporated into estimates of both biomass and $H R$. Our survey-based HR estimates for cod and whiting compared well with HR-converted fishing mortality ( $F$ ) estimates from analytical assessments for these two stocks. Of the non-target species tested, red gurnard (Chelidonichthys cuculus) recorded some annual HRs greater than those for cod or whiting; challenging "Pope's postulate" that Fon non-target stocks in an assemblage will not exceed that on target stocks. We relate $H R$ for each species to two corresponding maximum sustainable yield (MSY) reference levels; six non-target species (including three ray species) show annual $H R s \geq H R_{\text {MSY }}$. This result suggests that it may not be possible to conserve vulnerable non-target species when $F$ is coupled to that of target species. Based on biomass, $H R$, and $H R_{\text {MSY, }}$, we estimate "total allowable catch" for each non-target species.
Keywords: bycatch, catchability, data-poor stocks, elasmobranchs, Pope's postulate, stock assessment.

## Introduction

Bottom trawl surveys of the Celtic Sea shelf demersal assemblage record over 100 fish species. Commercial fisheries target predominately 10 of the largest and most abundant (Marine Institute, 2013), but survey catch per unit effort (cpue) indicates that several other species of little market value have similar overall abundance to these. Such abundant non-target species are frequently caught in commercial fishing gear (Pope et al., 2000), typically being discarded dead or dying (Benoit et al., 2013). It has proved difficult to quantify the effect of fishing pressure on non-target species, although long-term survey abundance trends in North Sea populations do imply fishing impact (Jennings et al., 1999). Fishing exerts mortality at some rate (the instantaneous death rate due to fishing $F$ ) on each species. Realized $F$ varies with behaviour and morphology, and can be expressed in terms of susceptibility and selectivity in the fishing gear, and the survival of fish after discarding (Zhou et al., 2010). Estimating $F$ for key target species is typically
undertaken using analytical stock assessments that are based on the age distribution of commercial catches. Such data are generally collected only for dominant target species and therefore such assessments are precluded as a means of estimating $F$ for non-target species.

In the current absence of empirical estimates of non-target $F$, some authors (e.g. Le Quesne and Jennings, 2012) suggest applying a precautionary principle such as "Pope's postulate" (Pope et al., 2000). Pope's postulate states that "fishing fleets generate a fishing mortality on non-target species which is less than or equal to that generated on the target species". It thus may provide upper limits on current mortality for "sensitive" species in a fishery and represents a useful management principle for data-poor situations (Jennings, 2013). This is a pragmatic approach, but may be too conservative in some fisheries and insufficiently precautionary in others. Only case-specific assessments can produce estimates of non-target mortality rate (Jennings, 2013). Nonetheless, accurate
estimates of fishing mortality on non-target species are required to fulfil international commitments on biodiversity conservation (e.g. CBD, 2010), and are central to the Ecosystem Approach to Fisheries Management (Garcia, 2003), which proposes to integrate fishing effects across exploited communities.

As an alternative to age-structured methods, Pope et al. (2000) proposed that the volume of catch (discards) in the fished component of a "local area" could be extrapolated to derive an estimate of $F$ for the whole of that local area. This method demands broad assumptions about catchability (the probability that an individual encountering fishing gear will be retained) and especially about spatial distribution of fish across the overall area. Where possible, a better approach may be to derive population biomass estimates from standardized fisheries-independent trawl surveys. The primary challenge here is that such surveys are not typically designed to estimate absolute fish population abundance, but to produce cpue indices of relative abundance. Many aspects of fishing gear selection and fish behaviour will affect catchability, and differences in catchability mean that surveys provide "biased perceptions of the actual abundance of different species and size classes at a particular time and location" (Fraser et al., 2007). Estimating absolute abundance from survey data requires a correction for catchability (ideally by species and length).

Survey catchability has been modelled (e.g. Harley and Myers, 2001), but Fraser et al. (2007) provide a list of empirically calculated length-based survey catchabilities for the most abundant NE Atlantic demersal fish species. Piet et al. (2009) developed a model that uses the catchability parameters from Fraser et al. (2007) to raise International Bottom Trawl Survey (IBTS) catches to spatial (ICES rectangle) estimates of biomass for the most abundant
species. The model simulates fishing of these species according to known fishing intensity and commercial gear selectivity, to derive model estimates of catch and then $F$. We benefit from access to data from an Irish monitoring programme that records discards of both commercial species and all non-target fish species. We use species catchabilities from Fraser et al. (2007) to derive population biomass estimates from survey data, and combine these with discard and landings records to yield empirical estimates of the proportion of biomass removed annually by fishing (harvesting rate $H R$ ) for 14 fish species in the Celtic Sea. To test the approach, our survey-based estimates of HR for "Celtic Sea" stocks of Atlantic cod (Gadus morhua) and Whiting (Merlangius merlangus) are compared with corresponding HRs converted from the $F$ estimates produced by analytical stock assessments. HRs for non-target species are then compared with HRs for the assessed stocks to test Pope's postulate.

Differences in life history mean that sustainable levels of $F$ vary considerably among demersal fish species. It is useful to be able to compare $F$ or $H R$ with appropriate reference levels for "sustainable" mortality. We present two estimates of sustainable $H R$ for each tested species, based on published approaches, and compare these estimated reference levels with observed annual $H R$ values. We then derive the "total allowable catch" (TAC) for each species that corresponds to a sustainable $H R$ given estimated biomass.

## Methods

Our study focuses on 15 International Council for the Exploration of the Seas (ICES) statistical rectangles in the Celtic Sea, with total area $\sim 52000 \mathrm{~km}^{2}$ (Figure 1). This area (approximating to ICES Area VIIg) was selected because it comprised the best overlap between available Irish Groundfish Survey (IGFS) data and Irish


Figure 1. Map of the Celtic Sea, showing ICES statistical rectangles included in the study.


Figure 2. Spatial distribution of Celtic Sea landings from the cod (left panel) and whiting (right panel) stocks (STECF, 2013) included in the study.
discard observer data, and captures much of the landings of the analytically assessed "Celtic Sea cod" stock (ICES, 2012a) and the "whiting in divisions VIIe-k" stock (ICES, 2012b; Figure 2). Each of these two stocks is assessed annually with an age-structured analytical assessment XSA (ICES, 2013a) to produce estimates of spawning-stock biomass (SSB), and importantly for this study, total-stock biomass (TSB) and $F$ at age. These assessments are primarily based on commercial landings data from the relevant stock area, but the most recent biomass estimates are adjusted ("tuned") using fisheries-independent survey data from the IGFS. In addition to the assessed cod and whiting stocks, we selected nine abundant non-target species from the IGFS, and also three exploited ray species of conservation interest. All selected species were deemed to be above the survey detection threshold, i.e. present in $>5 \%$ of all hauls and recording abundance $>5$ individuals $\mathrm{km}^{-2}$ (Trenkel and Cotter, 2009).

## Fish population biomass from survey data

The IGFS is a stratified random bottom-trawl survey that includes the Celtic Sea (Figure 1). This survey is operated in autumn (Q4) by the Irish Marine Institute using a GOV trawl fitted with a 20 mm codend liner. Standard International Bottom Trawl Survey (IBTS) protocol is followed. In a given year, trawl samples (designed to be 30 min duration at 4 knots) are collected at $1-10$ sites within each surveyed ICES rectangle. All fish captured in the IGFS are identified to species and measured (total length; $l_{\mathrm{cm}}$ ). For cod and whiting, and the selected non-target species, we calculate the annual biomass $B_{s, y}$ of species $s$ in the study area in year $y$, by summing over survey rectangles:

$$
\begin{equation*}
B_{s, y}=\sum_{k=1}^{K} B_{s, k, y}, \tag{1}
\end{equation*}
$$

where K is the number of rectangles in the study area and

$$
\begin{equation*}
B_{s, k, y}=A_{k} \bar{D}_{s, k, y} \tag{2}
\end{equation*}
$$

where $A_{k}$ is the sea surface area $\left(\mathrm{km}^{2}\right)$ of the $k$ th rectangle and $\bar{D}_{s, k, y}$ the mean biomass density $\left(\mathrm{kg} \mathrm{km}^{-2}\right)$ of species $s$ taken over $J_{k}$ individual survey trawls in rectangle $k$ and year $y$ :

$$
\begin{equation*}
\bar{D}_{s, k, y}=\frac{\sum_{j=1}^{J_{k}} D_{s, j, k, y}}{J_{k}} \tag{3}
\end{equation*}
$$

in which $D_{s, j, k, y}$ is the expanded biomass density of species $s$ estimated from trawl $j$, in survey rectangle $k$, and year $y$. Biomass of all species was expanded for survey trawlnet catchability, as follows:

$$
\begin{equation*}
D_{s, j, k, y}=\sum_{i=1}^{I}\left(D_{i, s, j, k, y} / q_{i, s}\right) \tag{4}
\end{equation*}
$$

in which $D_{i, s, j, k, y}$ is the observed biomass density of the $i$ th length class of species $s$ from trawl $j$, in survey rectangle $k$, and year $y$. The correction coefficient $q_{i, s}$ represents catchability and uses data from Fraser et al. (2007) resolved by species and length class.
$D_{s, j, k, y}$ for whiting only was calculated from doorspread [see Equation (7)] because this species is known to be herded into the path of the net by the sediment cloud stirred up by the otter doors and sweeps (e.g. Main and Sangster, 1981; Wardle, 1986). However, not all individual fish located between the doors are captured in the net. To account for this potential underestimate of biomass, for whiting only, we used:

$$
\begin{equation*}
D_{s, j, k, y}=\frac{\sum_{i=1}^{I}\left(D_{i, s, j, k, y} / q_{i, s}\right)}{h_{s}} \tag{5}
\end{equation*}
$$

where the coefficient $h_{s}$ is a length- and species-dependent herding factor, defined from Piet et al. (2009) as:

$$
h_{\text {whiting }}= \begin{cases}0.30 & \text { if } \quad l_{i}<29.5, \\ 0.75 & \text { if } \quad l_{i}<29.5,\end{cases}
$$

where $h$ is the probability that an individual fish located between the trawl doors will be herded in between the trawl wings and $l_{i}$ the median length of length class $i$.

Piet et al. (2009) assumed minimal herding for flatfish and rays, while the benthic habit and weak swimming characteristics of gurnards (Floeter and Temming, 2005) and dragonets (Takita et al., 1983) also suggest very little herding, and so $h$ was not applied for these species. However, some herding was assumed for the small roundfish Norway pout Trisopterus esmarkii and Poor cod Trisopterus minutus, meaning that wingspread biomass for these species [see Equation (7)] could be an overestimate. Observed
survey biomass for these species was thus expanded as:

$$
\begin{equation*}
D_{s, j, k, y}=h\left(\sum_{i=1}^{I}\left(\frac{D_{i, s, j, k, y}}{q_{i, s}}\right)\right) \tag{6}
\end{equation*}
$$

where the coefficient $h$ represents the probability that an individual of these two species that is swept by the trawl doorspread will enter the net: $h=R p_{\mathrm{n}}+(1-R) p_{\mathrm{d}}$, where $p_{\mathrm{n}}$ is the probability of entering the net if in front of the net wingspread and $p_{\mathrm{d}}$ is the probability of entering if in front of the doorspread but not wingspread, and $R$ is the ratio of wingspread to doorspread. Using $R=1 / 3$ and $p_{\mathrm{n}}=1$ and $p_{\mathrm{d}}=0.3, h=0.533$.

The observed biomass density estimate is obtained from the survey trawl swept-area as

$$
\begin{equation*}
=D_{i, s, j, k, y} \frac{w_{i, s, j, k, y}}{v_{j, k, y} \lambda_{j, k, y}} \tag{7}
\end{equation*}
$$

where $v_{j, k, y}$ is the length of the $j$ th trawl in rectangle $k$ and year $y$; $\lambda_{j, k, y}$ is that trawls wingspread (for all species except whiting, where it is that trawls doorspread-following Fraser et al., 2007), and $w_{i, s, j, k, y}$ is the total mass caught in that trawl in the $i$ th length class of species $s$. This mass is estimated from a length-weight relationship for each species as:

$$
\begin{equation*}
w_{i, j, k, y}=N_{i, j, k, y} \alpha l_{i}^{\beta} \tag{8}
\end{equation*}
$$

where $N_{i, j, k, y}$ is the number of individuals of length class $i$, caught in trawl $j$ in rectangle $k$ in year $y$, such that $l_{i}$ is the median length of length class $i$, and $\alpha$ and $\beta$ are species-specific values taken from Celtic Sea data for cod and whiting and from North Sea IBTS survey data for non-target species, which have insufficient weight records in the IGFS.

## Harvesting rate

## Discard data

Discard data came from an Irish observer programme that serves the Data Collection Regulation (EC No. 1639/2001). Fishing trips are sampled at a rate proportional to métier activity, with sampling coverage of the Irish fleet being $\sim 1 \%$ during the study period. Sampling trips are selected randomly, and so the distribution of fishing activity sampled is considered representative of the population as a whole (Marine Institute, 2011). Discard data were extracted by species, gear, quarter, and year. If a sampled fishing trip included hauls outside study rectangles, then the proportion of the fishing effort inside the area was used. Discard weight was raised to Irish fleet level by dividing it by the proportion of total Irish effort covered by discard sampling.

For cod and whiting, each of the Irish landings and discard values were raised to an international estimate by dividing by the annual proportion of the total catch landed by Irish vessels only in the stock area. This was considered a proxy for the proportion of total (international) effort (and hence discarding) accounted for by Irish vessels. For non-target species, discard records were raised according to the proportion (range $=51-58 \%$ in the study period) of annual international effort by mobile gears (kilowatt hours $=$ vessel engine power multiplied by time) in the study area recorded by Irish vessels (STECF, 2013). For years where effort for a given nation was not reported to STECF, the mean annual value for that nation was applied.

## Calculation

For each species $s$ in year $y$, a first $H R$ range for the study area was then estimated as:

$$
\begin{equation*}
H R_{s, y}^{B}=\frac{C_{s, y}}{C_{s, y}+\left(B_{s, y} \pm 1 \mathrm{s.d.}\right)} \tag{9}
\end{equation*}
$$

where $C_{s, y}$ is the total catch (landings and discards) of species $s$ in year $y$ and $B_{s, y}$ the expanded biomass (see above). $H R_{s, y}^{B}$ includes uncertainty in the estimate of survey biomass $B_{s, y}$. Confidence intervals around $B_{s, y}$ were estimated by repeated sub-sampling from the pool of observed hauls within year. To include (in addition) uncertainty in the estimation by the Irish observer scheme of discard rates, a second $H R$ range $H R_{s, y}^{B-\text { dis }}$ for the study area was then estimated as $H R_{s, y}^{B}$, but where $C_{s, y}=$ landings + discards $\times(1 \pm 20 \%)$; Rochet et al. (2002) estimated overall CV in discard estimates for the French fleet in the Celtic Sea as $20 \%$. $H R_{s, y}^{B-\text { dis }}$ may be considered most precautionary because the upper bound accounts for a scenario in which species biomass was overestimated by the survey, while the observer scheme underestimated discards (by $20 \%$, Rochet et al., 2002).

## Validation

As a validation exercise, $B_{s, y}$ and $H R_{s, y}^{B \text {-dis }}$ for cod and whiting were compared with TSB and $F$ values from respective stock assessments, where assessment $F$ s were converted to $H R[H R=1-\exp (-F)]$ for direct comparison. $B_{s, y}$ were smaller than assessment TSB for whiting, possibly reflecting the fact that the study covers only part of the assessed stock area. $H R_{\text {whiting, } y \text { B differed by an average of } 0.06}$ (12\%) from the mean annual assessment $H R$ estimates (Table 1). $B_{s, y}$ for cod were closer to assessment TSB, while $H R_{\text {cod, } y}^{B-\text { dis }}$ differed by an average of $0.05(16 \%)$ from assessment $H R$ with no years showing large discrepancy (Table 1). Similarity between surveybased and assessment HRs for both cod and whiting suggest that our approach generates estimates that can be reasonably applied to non-target species.

## Precautionary reference levels

To gain some insight into the likely ecological significance of observed $H R$ for non-target species, estimates for each species were compared with two sets of candidate reference levels:
(i) From a meta-analysis of 245 fish species, Zhou et al. (2012) suggested that $F_{\text {MSY }}$ could be estimated as 0.87 M for teleosts and $0.41 M$ for chondrichthyans. For most species, we calculated $H R_{\text {MSY }}$ reference points according to Zhou et al. (2012), but the value for witch (Glyptocephalus cynoglossus) was derived from ICES advice (ICES, 2013b). Estimates of natural mortality $M$ are from Gallagher et al. (2004) (rays) and FishBase (most other species). For red gurnard (Chelidonichthys cuculus), we used: $\ln (M)=1.44-0.982 \ln \left(t_{m}\right)$ (Hewitt and Hoenig, 2005), where $t_{m}$ is the maximum observed age reported in FishBase. An $H R$ limit reference point $\left(H R_{\text {lim }}\right)$ was also calculated by adding 1 s.d. to the $F_{\text {MSY: }}: M$ ratios of Zhou et al. (2012), i.e. $(0.41+0.09) M$ for elasmobranchs and $(0.87+0.05) M$ for teleosts, respectively.
(ii) For many of the demersal species in the Celtic Sea, Le Quesne and Jennings (2012) provide estimates of $F_{40}$ (the $F$ that reduces SSB-per-recruit to $40 \%$ of that in the absence of fishing). We used $F_{40}$ estimates from Table S1 in Le Quesne and Jennings (2012) to derive a list of $H R_{40}$ estimates.

Table 1. Survey-based $\left(B_{s, y}\right)$ and assessment (TSB) estimates of total biomass and harvesting rate for cod and whiting.

|  | Whiting |  |  |  | Atlantic cod |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 2008 | 2009 | 2010 | 2011 | 2008 | 2009 | 2010 | 2011 |
| International landings | 9093 | 6382 | 8563 | 9639 | 3639 | 3263 | 3229 | 4737 |
| International discards | 3479 | 7569 | 4184 | 3188 | 1487 | 1351 | 1833 | 7541 |
| $B_{s, y} \pm 1$ s.d. in study area | $8385 \pm 871$ | $17151 \pm 905$ | $12761 \pm 635$ | $19674 \pm 1198$ | $5928 \pm 399$ | $3638 \pm 386$ | $14113 \pm 1016$ | $27293 \pm 2462$ |
| Assessment TSB | 29151 | 44568 | 61056 | 61623 | 9216 | 9781 | 23145 | 23358 |
| $H R^{B}$ | 0.58-0.63 | 0.44-0.46 | 0.49-0.51 | 0.38-0.41 | 0.45-0.48 | 0.53-0.59 | 0.25-0.28 | 0.29-0.33 |
| $H R^{B \_ \text {dis }}$ | 0.57-0.63 | 0.42-0.48 | 0.48-0.52 | 0.37-0.42 | 0.44-0.48 | 0.53-0.59 | 0.24-0.29 | 0.28-0.34 |
| Assessment HR | 0.51 | 0.46 | 0.44 | 0.32 | 0.51 | 0.52 | 0.38 | 0.31 |

$H R_{s, y}^{B}$ is a range incorporating uncertainty in survey biomass estimates $\left(H R_{s, y}\right.$ at $\left.B_{s, y} \pm 1 s . d.\right)$. $H R_{s, y}^{B \text { dis }}$ is a range incorporating uncertainty in estimates of both survey biomass $B_{s, y}$ and discards. Assessment $H R=1-\exp (-F)$, using ages 2-5 for cod and 2-7 for whiting (ICES, 2013a). International landings and discard estimates are also shown. All weights are tonnes.

Table 2. Survey-based (IGFS) estimates of total-stock biomass in tonnes ( $B_{s, y} \pm 1$ s.d.), 95 th percentile of length $\left(I_{95}, \mathrm{~cm}\right)$ from the IGFS data, and natural mortality $(M)$ from Fishbase for non-target demersal fish species in the study area of the Celtic Sea.

| Common name | Latin name | $L_{95}$ | M | $\begin{aligned} & 2008 \\ & B \end{aligned}$ | $\begin{aligned} & 2009 \\ & B \end{aligned}$ | $\begin{aligned} & 2010 \\ & B \end{aligned}$ | $\begin{aligned} & 2011 \\ & \text { B } \end{aligned}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Lesser spotted dogfish | Scyliorhinus canicula | 67 | 0.37 | $32272 \pm 8618$ | $25771 \pm 1326$ | $42734 \pm 2927$ | $46734 \pm 4996$ |
| Spotted ray | Raja montagui | 61 | 0.41 | $2832 \pm 336$ | $7714 \pm 630$ | $13652 \pm 666$ | $9728 \pm 1195$ |
| Thornback ray | Raja clavata | 78 | 0.20 | NA | $3709 \pm 320$ | $6136 \pm 659$ | $2113 \pm 234$ |
| Cuckoo ray | Leucoraja naevus | 63 | 0.37 | $4506 \pm 653$ | $2766 \pm 326$ | NA | $5004 \pm 453$ |
| Witch | G. cynoglossus | 40 | 0.18 | $7070 \pm 713$ | $31663 \pm 4299$ | $6920 \pm 235$ | $12951 \pm 391$ |
| Dab | Limanda limanda | 26 | 0.26 | $1040 \pm 147$ | $2653 \pm 270$ | $2084 \pm 140$ | $4179 \pm 508$ |
| Long rough dab | H. platessoides | 22 | 0.32 | $1299 \pm 89$ | $1208 \pm 80$ | $1610 \pm 66$ | $1539 \pm 78$ |
| Poor cod | Trisopterus minutus | 21 | 1.00 | $83570 \pm 6797$ | $79816 \pm 2959$ | $48918 \pm 2665$ | $39395 \pm 2157$ |
| Grey gurnard | Eutrigla gurnardus | 31 | 0.29 | $3213 \pm 279$ | $3317 \pm 328$ | $10261 \pm 756$ | $13015 \pm 1144$ |
| Red gurnard | C. cuculus | 32 | 0.20 | $280 \pm 36$ | $385 \pm 38$ | $484 \pm 24$ | $398 \pm 39$ |
| Dragonets | Callionymus spp. | 25 | 0.91 | $1041 \pm 135$ | $1338 \pm 137$ | $2113 \pm 79$ | NA |
| Norway pout | Trisopterus esmarkii | 21 | 1.54 | $16855 \pm 1058$ | $43256 \pm 2138$ | $75228 \pm 6294$ | $104805 \pm 18867$ |

A precautionary mean total allowable catch (TAC) or total allowable discards (TAD) for each species was estimated by averaging the lower bounds of $B_{s, y}$ and then multiplying by the most conservative (lowest) HR reference level.

## Survey trends

Time-series of cpue $\left(\mathrm{kg} \mathrm{km}^{-2}\right)$ in the IGFS and also the UK West Coast Groundfish Survey (WCGFS) were used as a visual descriptor of changes in species relative abundance that might partly reflect $F$. The WCGFS was discontinued in 2004 but has some data for the study region extending to 1986 and is assumed to describe fish abundance at an earlier stage of exploitation history (Shephard et al., 2011).

## Results

## Survey biomass

$B_{s, y}$ varied among species as expected, with low estimates for "k-strategy" (low M) species such as the rays and higher estimates for "r-strategy" (high $M$ ) species such as Norway pout and Poor cod (Table 2).

## Harvesting rate

Non-target species showed a wide range of $H R_{s, y}^{B-}$ dis, with some missing values due to incomplete discard data (Table 3). Two roundfish (red gurnard and grey gurnard Eutrigla gurnardus) recorded some $H R_{s, y}^{B-\text { dis }}$ in the same range as estimated for cod, with some values for red gurnard exceeding the maxima recorded for assessed stocks of both cod and whiting (Table 3).

## Reference levels

Of the teleost species, dab (Limanda limanda), red gurnard, and grey gurnard showed at least one annual $H R$ greater than one of the proposed $H R$ reference levels, while some $H R s$ for red gurnard were greater than both reference levels. All tested elasmobranchs recorded at least one $H R_{s, y}^{B-d i s}$ with upper bounds greater than or equal to one of the tested reference levels. Spotted ray Raja montagui, which has the lowest $l_{95}$ and greatest $M$ among the elasmobranchs, and might thus be expected to be the most resilient, recorded lowest $H R_{s, y}^{B-d i s}$ in this group (Table 3). The estimated TACs varied strongly among species, with red gurnard TAC estimated at only 56 t while Norway pout TAC was $>20000 \mathrm{t}$.

## Discussion

The ecosystem approach to fisheries requires information on population size and fishing mortality of non-target species within exploited fish assemblages. We use catchability-expanded survey records and catch (landings + discards) data to estimate biomass and $H R$ for two assessed and several non-target demersal fish species. $H R_{\text {cod, } y}^{B-\text { dis }}$ and $H R_{\text {whiting,y }}^{B-\text { dis }}$ are close to analytical stock assessment estimates for these species, suggesting that our method may produce reasonable estimates for the non-target species. We aim to present a pragmatic method that can yield information on the state of surveyed but unassessed stocks. This requires certain assumptions that we discuss with reference to the fisheries survey criteria presented by Trenkel and Cotter (2009).
Table 3. Harvesting rate estimates for non-target demersal fish in the study area of the Celtic Sea.


## Survey data

## Study area vs. fish "stock" distributions

To validate our approach, we needed to make a direct comparison between survey-based estimates of biomass and mortality for given stocks, and corresponding estimates from analytical assessments. We identified an area of the Celtic Sea (Figure 2) corresponding to the core area (greatest landings: STECF, 2013) of assessed cod and whiting stocks, and for which survey and catch data were available. Comparison of survey-based and assessment biomass estimates suggested that we probably captured much of the cod population but only a component of the whiting population-this was anticipated from the known range of these stocks. Comparison of survey and assessment HRs suggest that the study area allowed quite accurate estimates of fishing mortality for both cod and whiting. For simplicity in continuing the current "worked example" of our approach, we used the same study area for non-target "stocks". Our results thus represent biomass and $H R$ estimates for that component of each non-target stock within our study area in a given year. When applying the method at, for example, MSFD Subregion scale, there may be ecological justification for defining species-specific ranges from survey cpue data. A potential difficulty is that these ranges are likely to cross national survey and discard sampling boundaries, and so synthesis of disparate dataseries may be required.

In our biomass estimates $B_{s, y}$, we stratify the survey data by ICES rectangle, as in the analytical stock assessments. This simple approach incorporates spatial heterogeneity in population density without demanding complex evaluation of density patterns and temporal changes in these patterns. An alternative approach might be to stratify the survey data in a more dynamic way, based on observed spatial density of fish within and among years.

## Catchability

We applied $q_{i, s}$ from Fraser et al. (2007) to raise survey cpue by species and length class to total biomass within the study area. Fraser et al. (2007) calculated catchability parameters for assessed species by quantifying the ratios between catch at length in North Sea survey data and numbers-at-age (converted to length) in analytical stock assessments for the same stocks. For whiting, we follow Piet et al. (2009) in using the relevant Fraser et al. (2007) catchability parameters with an additional trawl door herding factor $h$. Fraser et al. (2007) estimated catchabilities for non-target species by comparing catch between beam trawl and GOV hauls undertaken in "approximately the same time and place". This method assumes that $q_{i, s}$ in the beam trawl are fairly stable relative to the GOV, and close to 1 . We suggest that this is a reasonable assumption for flatfish and rays (Piet et al., 2009), gurnards (Floeter and Temming, 2005), and dragonets (Takita et al., 1983), which are unlikely to have a very active or sustained escape response to fishing gear. Many other small non-target demersal species also show benthic behaviour, and probably low herding response. For Poor cod and Norway pout, we assume more active herding and escape behaviour and apply a similar size-based herding factor $h$ (Piet et al., 2009).

Fraser et al. (2007) derived $q_{i, s}$ for the standard GOV trawl used in the Q3 North Sea IBTS. However, the same parameters have been widely applied to the separate Q1 North Sea IBTS (e.g. Collie et al., 2013; Heath et al., 2013) and result in very similar species biomass trends between surveys (Greenstreet et al., in preparation). We thus assume that Fraser et al. (2007) can be used as a first-order approach to raise catch in similar survey GOV trawls in different areas/seasons. This assumption is upheld by similarity between
survey-based and assessment $H R$ estimates for cod and whiting in our study area.

Catchability for given species can vary with size, and this is already incorporated into the Fraser et al. (2007) catchability parameters, which typically comprise a catchability curve with length. Catchability $q_{i, s}$ may also vary with season due to changes in spatial distribution of stock components (size classes) and in their availability to the survey gear. Without species-level information on such effects in the study area, it is difficult to evaluate whether the Q4 IGFS undersamples certain population components. Trenkel and Cotter (2009) suggest that if species cpue trends are similar between different surveys in a given area, then $q_{i, s}$ may be similar between surveys. This is the case between the Q1 and Q3 IBTS in the North Sea (Greenstreet et al., in preparation).

## Harvesting rate

Our estimates of $H R$ depend on survey, observer, and commercial landings data. We have to assume that the landings data are accurate, and so the two major potential sources of uncertainty in our approach are the survey-based estimates of species biomass (which include uncertainty in catchability $q_{i, s}$ ) and the observer estimates of discarding. We attempt to address both these sources of uncertainty, although uncertainty in $q$ is not explicitly addressed. The survey follows a robust scientific protocol in which samples are located randomly within ICES rectangles. By subsampling (random $10 \%$ deletion) from the pool of survey hauls in each year, we address the potential effect of haul location and inter-haul variation on annual species biomass estimates, while maintaining spatial stratification by ICES rectangle. The discard observer scheme is more ad hoc, with overall fleet coverage stratified by métier, but actual sampling depending partly on access to fishing vessels, weather, seasonality in fishing patterns, etc. We thus take a more flexible approach to uncertainty in the discard data, incorporating into $H R$ estimates the effect of $\pm 20 \%$ error (Rochet et al., 2002) in the estimate of annual discards of a given species. As expected, the range in annual $H R$ for each species tends to be wider when both sources of uncertainty are included (Table3). The most precautionary application of $H R$ for management would be to use the upper estimate, and this is the approach we take when proposing TACs.

## Validation

We "validate" our method by comparing $B_{s, y}$ and $H R_{s, y}^{B-\text { dis }}$ for cod and whiting to estimates from corresponding analytical stock assessments. We note that while the stock assessments are predominately based on age-structured landings data, they use the IGFS to tune (adjust) biomass series. This means that the two approaches to estimating biomass and mortality are not completely independent. Survey cpue generally has most influence on terminal estimates of fishing mortality and SSB in the assessments, and so the most recent 3 (approximately) years, assessment $F$ and SSB are most strongly influenced by the survey. For earlier years, the assessments tend to converge on the catch data according to the traditional convergence properties of VPA-based assessments. On this basis, the IGFS will not strongly influence cod and whiting assessment outputs (which we took from 2013 ICES WG reports) for the earlier years used in our study (2008-2010), but there will be some non-independence for the final year (2011). Even for 2011, we suggest that possible nonindependence does not matter, since our intention is to provide survey-based estimates that can serve, for unassessed populations, the same purpose as age-structured assessments. Correspondingly, we find that we can use survey data directly and derive results that
are similar to those produced by those assessments. This result suggests that for other species, survey data may also be sufficient to produce estimates that would be similar to those from age-structured assessment if such assessments were conducted.

## Reference levels

Fishing mortality reference points provide insight into the likely ecological significance of observed $H R s$ for non-target species. Slower growing, later maturing, and less fecund species (e.g. many elasmobranchs) are expected to experience greater negative impact for a given $H R$ (e.g. Jennings et al., 1999; Garcia et al., 2008; Le Quesne and Jennings, 2012). High levels of $F$ for the ray species probably reflect strong susceptibility to capture in mobile fishing gears (Cedrola et al., 2005). Patches of low fishing effort in the Celtic Sea can act as de facto refuges for some elasmobranchs (Shephard et al., 2012), but our HR estimates suggest that additional conservation measures are probably necessary to reach precautionary reference levels for $F$ in these species.

The high HRs for both gurnard species are notable (Table 3). Rochet et al. (2002) found that red and grey gurnards (and boarfish Caprosaper) were the most common non-target species discarded in the Celtic Sea; discarding of the highly perishable red gurnard increased with trip length. This high level of discarding for red gurnard, compared with our small abundance estimates for this species (Table 2), helps explain observed high HR. Gurnards may be particularly susceptible to capture in trawl gear because of morphology (broad square head and long spiny fins) and behaviour (sit-and-wait or stalk predation with little muscle capacity for sustained escape swimming; Floeter and Temming, 2005).

We estimate a mean TAC (2008-2011) for each study non-target species. These TACs currently refer only to the study area (approximately ICES VIIg), but the method presented may have considerable utility in the context of the upcoming EU landing obligation (EU COM/2013/0889 final-2013/0436). Understanding fishing mortality of vulnerable species and being able to relate this to a specific volume of landings will enable managers to make informed decisions about conservation of these species. A significant problem in this regard is that $F$ of vulnerable species is closely tied to that of target species, and reducing target $F$ to a level consistent with conservation of the most vulnerable species may restrict fishing activities to a socially unacceptable degree. A potential solution might be the use of spatial management to de-couple target and non-target $F$, possibly with particular application to elasmobranchs (e.g. Shephard et al., 2012).

## Survey trends

The IGFS time-series showed little trend in standardized abundances (haul density) of non-target species. The WCGFS recorded visible declines in several species that we find to have high $H R s$, including red gurnard and cuckoo ray (Figure 3). These data suggest that non-target species experiencing high fishing mortality can show decadal population declines. Spotted ray, for which we observed $H R>H R_{\text {msy }}$ in 2008 only, also demonstrated a declining trend in the IGFS

## Conclusions

$H R$ can be estimated for surveyed non-target species, and related to precautionary reference levels. At least two non-target species discarded in the Celtic Sea experience $H R$ s greater than those recorded for assessed cod and whiting stocks, meaning that Pope's postulate does not hold in the study area. Some vulnerable non-target species are "exploited" at rates greater than precautionary reference levels,


Figure 3. Standardized biomass (mean haul density, $\mathrm{kg} \mathrm{km}^{-2}$ ) for cod, whiting, and 12 non-target species in the UK Celtic Sea West Coast Groundfish Survey (WCGFS).
and some of those show declines in survey cpue. It is possible to calculate the catch (TAC or TAD) associated with precautionary reference levels and thus monitor exploitation of vulnerable species.

## Acknowledgements

The research leading to these results has received funding from the European Community's Seventh Framework Programme (FP7/ 2007-2013) under grant agreement MYFISH number 289257. DGR and KDF also acknowledge funding from a Beaufort Marine Research Award, carried out under the Sea Change Strategy and the Strategy for Science Technology and Innovation (2006-2013), with the support of the Marine Institute, funded under the Marine Research Subprogramme of the National Development Plan 2007-2013.

## References

Benoît, H. P., Plante, S., Kroiz, M., and Hurlbut, T. 2013. A comparative analysis of marine fish species susceptibilities to discard mortality: effects of environmental factors, individual traits, and phylogeny. ICES Journal of Marine Science, 70: 99-113.
CBD. 2010. X/2. The Strategic Plan for Biodiversity 2011-2020 and the Aichi Biodiversity Targets. Convention on Biological Diversity, Nagoya, Japan. 18-29 October 2010. UNEP/CBD/COP/DEC/X/ 2. 13 pp .

Cedrola, P. V., González, A. M., and Pettovello, A. D. 2005. Bycatch of skates (Elasmobranchii: Arhynchobatidae, Rajidae) in the Patagonian red shrimp fishery. Fisheries Research, 71: 141-150.
Collie, J., Rochet, M-J., and Bell, R. 2013. Rebuilding fish communities: the ghost of fisheries past and the virtue of patience. Ecological Applications, 23: 374-391.
Floeter, J., and Temming, A. 2005. Analysis of prey size preference of North Sea whiting, saithe, and grey gurnard. ICES Journal of Marine Science, 62: 897-907.
Fraser, H. M., Greenstreet, S. P., and Piet, G. J. 2007. Taking account of catchability in groundfish survey trawls: implications for estimating demersal fish biomass. ICES Journal of Marine Science, 64: 1800-1819.
Gallagher, M. J., Nolan, C. P., and Jeal, F. 2004. An investigation of the Irish ray fishery in ICES divisions VIIa and VIIg. Journal of Northwest Atlantic Fisheries Science, 35: 1-13.
Garcia, S. M. (Ed.). 2003. The ecosystem approach to fisheries: issues, terminology, principles, institutional foundations, implementation and outlook (No. 443). Food \& Agriculture Organization of the United Nations.
García, V. B., Lucifora, L. O., and Myers, R. A. 2008. The importance of habitat and life history to extinction risk in sharks, skates, rays and chimaeras. Proceedings of the Royal Society London, Series B, 275: 83-89.
Harley, S. J., and Myers, R. A. 2001. Hierarchical Bayesian models of length-specific catchability of research trawl surveys. Canadian Journal of Fisheries and Aquatic Sciences, 58: 1569-1584.
Heath, M. R., Culling, M. A., Crozier, W. W., Fox, C. J., Gurney, W. S., Hutchinson, W. F., and Carvalho, G. R. 2013. Combination of genetics and spatial modelling highlights the sensitivity of cod (Gadus morhua) population diversity in the North Sea to distributions of fishing. ICES Journal of Marine Science, 71: 794-807.
Hewitt, D. A., and Hoenig, J. M. 2005. Comparison of two approaches for estimating natural mortality based on longevity. Fishery Bulletin, 103: 433-437.
ICES. 2012a. Celtic Sea and West of Scotland Cod in Divisions VIIe-k (Celtic Sea cod).ICES Advice Document. http://www.ices.dk/sites/ pub/Publication\%20Reports/Advice/2012/2012/cod-7e-k.pdf.
ICES. 2012b. Celtic Sea and West of Scotland Whiting in Divisions VIIe-k. ICES Advice Document. http://www.ices.dk/sites/pub/ Publication\%20Reports/Advice/2012/2012/whg-7e-k.pdf.

ICES. 2013a. Report of the Working Group for Celtic Seas Ecoregion (WGCSE), 8-17 May 2013, Copenhagen, Denmark. ICES CM 2013/ACOM: 12. 1986 pp. http://www.ices.dk/sites/pub/ Publication\%20Reports/Expert\%20Group\%20Report/acom/2013/ WGCSE/wgcse_2013.pdf.
ICES. 2013b. Witch in Subarea IV and Divisions IIIa and VIId. ICES Advice Document. http://www.ices.dk/sites/pub/Publication\  Reports/Advice/2013/2013/wit-nsea.pdf.
Jennings, S., Greenstreet, S. P. R., and Reynolds, J. D. 1999. Structural change in an exploited fish community: a consequence of differential fishing effects on species with contrasting life histories. Journal of Animal Ecology, 68: 617-627.
Jennings, S. 2013. When can "principles" support advice on fisheries and environmental management? ICES Journal of Marine Science, 70: 726-733.
Le Quesne, W. J., and Jennings, S. 2012. Predicting species vulnerability with minimal data to support rapid risk assessment of fishing impacts on biodiversity. Journal of Applied Ecology, 49: 20-28.
Main, J., and Sangster, G. I. 1981. A study of the sand clouds produced by trawl boards and their possible effect on fish capture. Department of Agriculture and Fisheries for Scotland. 20 pp.
Marine Institute. 2011. Atlas of Demersal Discarding, Scientific Observations and Potential Solutions, Marine Institute, Bord Iascaigh Mhara, September 2011. ISBN 978-1-902895-50-5. 82 pp .
Marine Institute. 2013. The Stock Book: Report to the Minister for Agriculture, Food and the Marine. Annual 2013 Review of Fish Stocks with Management Advice for 2014. Irish Marine Institute. 534 pp .
Piet, G. J., Van Hal, R., and Greenstreet, S. P. R. 2009. Modelling the direct impact of bottom trawling on the North Sea fish community to derive estimates of fishing mortality for non-target fish species. ICES Journal of Marine Science, 66: 1985-1998.
Pope, J. G., Macdonald, D. S., Daan, N., Reynolds, J. D., and Jennings, S. 2000. Gauging the impact of fishing mortality on non-target species. ICES Journal of Marine Science, 57: 689-696.
Rochet, M. J., Péronnet, I., and Trenkel, V. M. 2002. An analysis of discards from the French trawler fleet in the Celtic Sea. ICES Journal of Marine Science, 59: 538-552.
Shephard, S., Gerritsen, H. D., Kaiser, M. J., and Reid, D. G. 2012. Spatial heterogeneity in fishing creates de facto refugia for endangered Celtic Sea elasmobranchs. Plos One, doi:10.1371/journal. pone. 0049307.
Shephard, S., Greenstreet, S. P. R., and Reid, D. G. 2011. Interpreting the large fish indicator for the Celtic Sea. ICES Journal of Marine Science, 68: 1963-1972.
STECF. 2013. Scientific, Technical and Economic Committee for Fisheries. http://stecf.jrc.ec.europa.eu/web/stecf/ewg1313. Appendix 05 . Effective effort by rectangle by country.
Takita, T., Iwamoto, T., Kai, S., and Sogabe, I. 1983. Maturation and spawning of the dragonet, Callionymus enneactis, in an aquarium. Japanese Journal of Ichthyology, 30: 221-226.
Trenkel, V. M., and Cotter, J. 2009. Choosing survey time series for populations as part of an ecosystem approach to fishery management. Aquatic Living Resources, 22: 121-126.
Wardle, C. S. 1986. Fish behaviour and fishing gear. In The Behaviour of Teleost Fishes, pp. 463-495. Springer US, New York.
Zhou, S., Smith, A. D., Punt, A. E., Richardson, A. J., Gibbs, M., Fulton, E. A., Pascoe, S., et al. 2010. Ecosystem-based fisheries management requires a change to the selective fishing philosophy. Proceedings of the National Academy of Science of the United States of America, 107: 9485-9489.
Zhou, S., Yin, S., Thorson, J. T., Smith, A. D., and Fuller, M. 2012. Linking fishing mortality reference points to life history traits: an empirical study. Canadian Journal of Fisheries and Aquatic Sciences, 69: 1292-1301.

