# Calibration

2023-08-09

#### data

Generate covariate, treatment, outcome data-structure

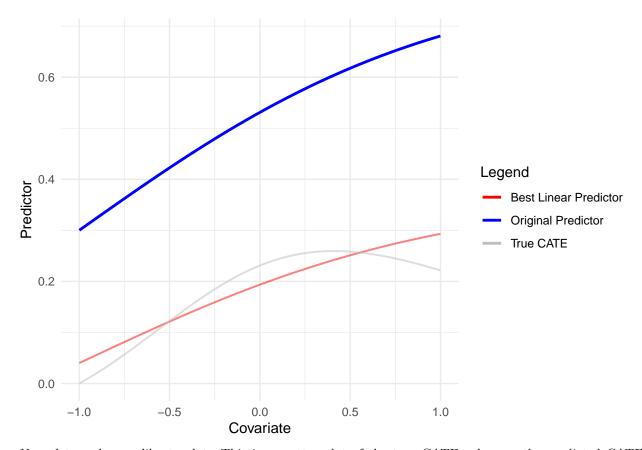
```
set.seed(12345)
n <- 2000
# covariate
W \leftarrow runif(n, -1, 1)
# treatment
pi <- plogis(0.5*W)</pre>
A <- rbinom(n, size = 1, pi)
# outcome regression and cate
mu0 <- plogis(W)</pre>
mu1 <- plogis(1 + 2*W)</pre>
cate <- mu1 - mu0
# potential outcomes
YO <- rbinom(n, size = 1, mu0)
Y1 <- rbinom(n, size = 1, plogis(1 + 2*W))
# observed outcome
Y <- ifelse(A==1, Y1, Y0)
```

train initial predictor and make calibration plot (predictor vs CATE)

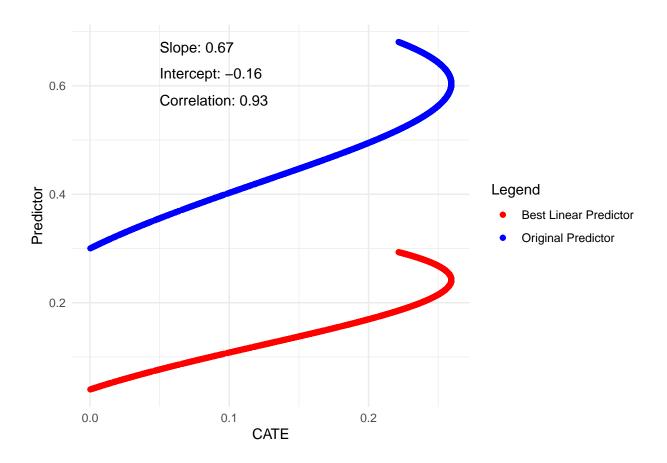
```
# use machine learning to obtain initial predictor of ITE Y_1 - Y_0
# for simplicity, we define our predictor tau.hat as a fixed function.
tau.hat <- plogis(1 + W ) - 0.2
```

Lets plot the predictor, the BLP, and the true CATE as a function of the covariate.

```
## Warning: Using 'size' aesthetic for lines was deprecated in ggplot2 3.4.0.
## i Please use 'linewidth' instead.
## This warning is displayed once every 8 hours.
## Call 'lifecycle::last_lifecycle_warnings()' to see where this warning was
## generated.
```



Now, lets make a calibrate plot. This is a scatter plot of the true  $\operatorname{CATE}$  values vs the predicted  $\operatorname{CATE}$  values.



## Linear calibration with Victor's BLP method

Now lets use Victor's BLP method to linearly calibrate our original predictor.

```
# unbiased surrogate outcome for CATE/ITE
pseudo_outcome <- cate + (A/pi) * (Y - mu1) - ((1-A)/(1-pi)) * (Y - mu0)

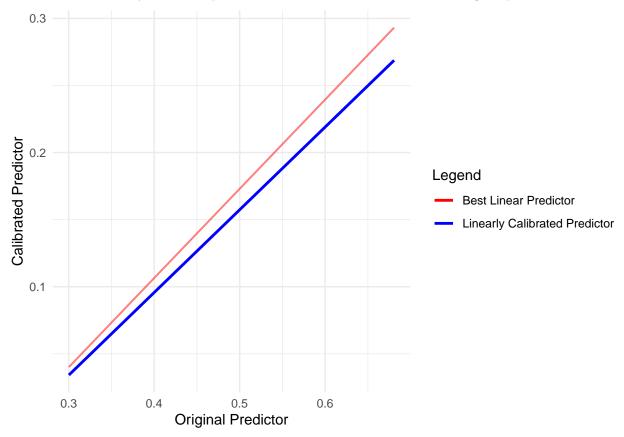
# fit best linear predictor of tau.hat of the surrogate outcome
# provides estimat of BLP of ITE/CATE
fit <- lm(pseudo_outcome ~ tau.hat, data = data.frame(tau.hat, pseudo_outcome))
intercept <- coef(fit)[1]
slope <- coef(fit)[2]

# get linear calibrated predictor, i.e. BLP given tau.hat
tau.BLP.hat <- intercept + slope * tau.hat

cor_tau <- cor(tau.BLP.hat, cate)
# Calculate the regression coefficients
fit <- lm(cate ~ tau.hat, data = data.frame(tau.hat, cate))
intercept <- coef(fit)[1]
slope <- coef(fit)[2]</pre>
```

tau.BLP.oracle <- intercept + tau.hat \* slope</pre>

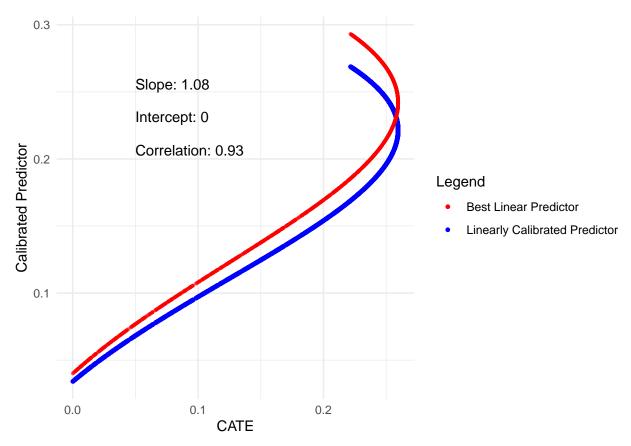
We see that the linearly calibrated predictor is a linear transformation of the original predictor.



### Check calibration of BLP-corrected predictor

Lets check the new calibration plot of the linearly calibrated predictor.

The following plot shows how the treatment effect predictions assigned to individuals (y-axis) varies as a function of the actual conditional average treatment effect of the individuals.



We see now that the linearly calibrated predictor is well calibrated in a linear sense. The fit cannot be improved by fitting a linear model on top of the predictor. We have correctly estimated the BLP.

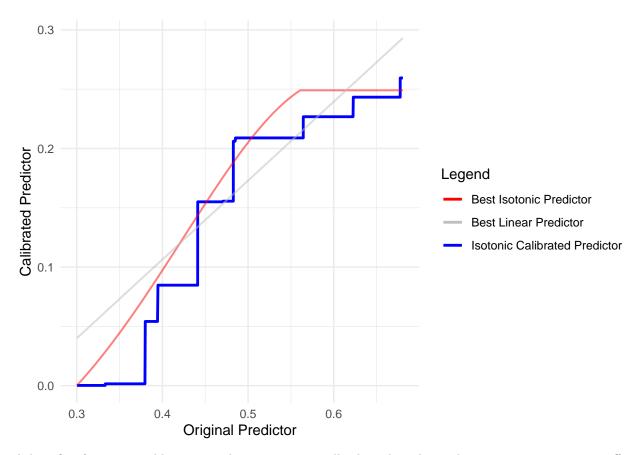
However, we see that the BLP approach is unable to correct for the poor calibration (flattening) at the end regions. This is because BLP is parametric and can only calibrate by applying linear transformations to the predictor. A more nonparametric approach can correct for nonlinear transformation.

#### Causal isotonic calibration

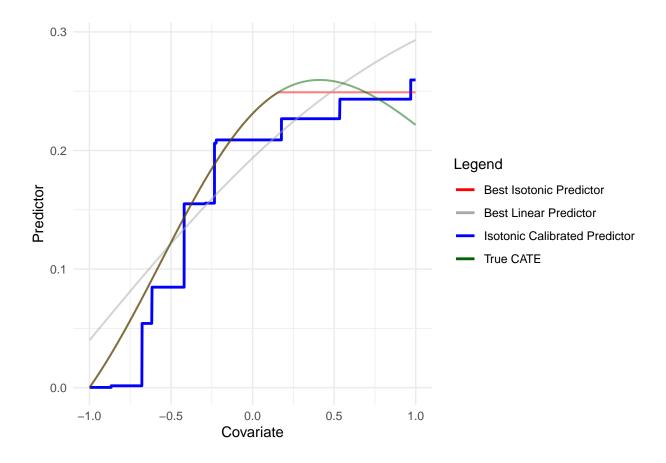
Now lets use causal isotonic calibration to nonparametrically calibrate our original predictor. Causal isotonic calibration is able to correct for any monotone transformations of the original predictor. Since linear transformations (with positive slope) are monotone, this method is strictly more powerful than the BLP approach.

```
# unbiased surrogate outcome for CATE/ITE
pseudo_outcome <- cate + (A/pi) * (Y - mu1) - ((1-A)/(1-pi)) * (Y - mu0)
# fit isotonic regression of pseudo outcome
tau.iso.hat <- as.stepfun(isoreg(tau.hat, pseudo_outcome))(tau.hat)
# note isotonic regression overfits at very end boundaries of tau.hat
# tmp fix: bound predictions into range of cate.
tau.iso.hat <- pmin(pmax(tau.iso.hat, min(cate)), max(cate))</pre>
```

We see that the isotonic calibrated predictor is a monotone piece-wise constant transformation of the original predictor.



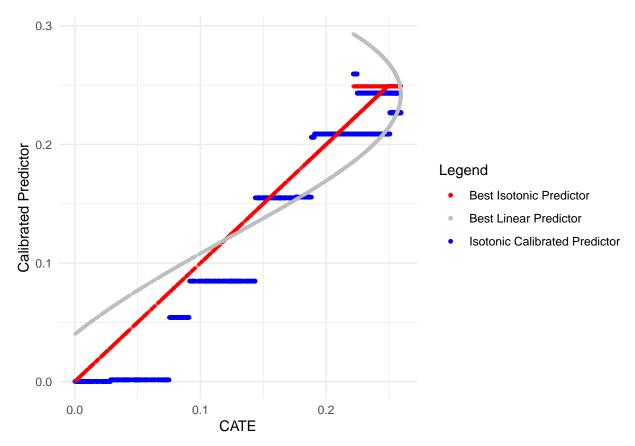
A benefit of isotonic calibration is that it automatically does data-driven heterogeneous treatment effect subgroup identification. The piece-wise constant values of the calibrated predictor defines subgroups (or bins) of individuals. Individuals with a given bin have conditional treatment effects that are quantitatively similar (given only information provided by the original predictor). Moreover, the conditional treatment effect of a given subgroup is meaningfully different form those of other subgroups.



# ${\bf Check\ calibration\ of\ isotonic\text{-}corrected\ predictor}$

Lets check the new calibration plot of the isotonic calibrated predictor.

The following plot shows how the treatment effect predictions assigned to individuals (y-axis) varies as a function of the actual conditional average treatment effect of the individuals.



We see now that the linearly calibrated predictor is well calibrated in a linear sense. The fit cannot be improved by fitting a linear model on top of the predictor. We have correctly estimated the BLP.

However, we see that the BLP approach is unable to correct for the poor calibration (flattening) at the end regions. This is because BLP is parametric and can only calibrate by applying linear transformations to the predictor. A more nonparametric approach can correct for nonlinear transformation.