DGZ Notebook

Data cleaning for DGZ

Load the libraries

Function to apply SHA-256 hashing

```
# Function to apply SHA-256 hashing
sha256_hash <- function(data) {
  openss1::sha256(data)
}</pre>
```

Data loading

```
barometer_dt_raw <- readxl::read_excel("../Data/DGZ/DECIDE_MTA_UGENT_14nov2022.xlsx")
barometer_aero_cult_raw <- readxl::read_excel("../Data/DGZ/DECIDE_MTA_UGENT_BAC_AERO_14nov2022.xlsx")
barometer_myco_cult_raw <- readxl::read_excel("../Data/DGZ/DECIDE_MTA_UGENTBAC_MYCO_14nov2022.xlsx")</pre>
```

Data manipulation AEROBIC CULTURE results

```
barometer_aero_cult <- barometer_aero_cult_raw %>%
  dplyr::rename(
   Filenumber = Dossiernummer,
   Pathogen_identification = 'KIEMSTAAL IDENTIFICATIE',
   Pathogen_result = 'KIEMSTAAL RESULTAAT',
   Samplenumber = 'Staalnummer'
 ) %>%
  dplyr::mutate(
   Parameter_code = 'BAC_AERO',
   Result = 'OK'
  ) %>%
  dplyr::select(
   Filenumber,
   Pathogen identification,
   Pathogen_result,
   Parameter_code,
```

```
Samplenumber,
    Result
  ) %>%
  dplyr::filter(
    Pathogen identification %in% c("Pasteurella multocida", "Mannheimia haemolytica", "Histophilus somni"
  dplyr::distinct() %>%
  dplyr::mutate(
    Filenumber_anon = sha256_hash(as.character(Filenumber)),
    Samplenumber_anon = sha256_hash(as.character(Samplenumber))
  dplyr::select(-Filenumber, -Samplenumber)
Intermediate table is needed
df_samples <- data.frame(</pre>
  Result = c('OK', 'OK', 'OK', 'OK'),
  Parameter_code = c('BAC_AERO','BAC_AERO','BAC_AERO', 'BAC_MYCOPLASMA'),
  Diagnostic_test= c('Culture','Culture','Culture', 'Culture'),
  Pathogen_identification=c("Pasteurella multocida", "Mannheimia haemolytica", "Histophilus somni", 'Myco
```

Data manipulation MYCOPLASMA CULTURE results

```
# Data manipulation MYCOPLASMA CULTURE results
barometer_myco_cult <- barometer_myco_cult_raw %>%
  dplyr::rename(
   Filenumber = Dossiernummer,
   Pathogen_identification = 'KIEMSTAAL IDENTIFICATIE',
   Mycoplasma result = 'KIEMSTAAL RESULTAAT',
   Samplenumber = 'Staalnummer'
 ) %>%
  dplyr::mutate(
   Parameter_code = 'BAC_MYCOPLASMA',
   Result = 'OK'
  ) %>%
  dplyr::select(
   Filenumber,
   Pathogen_identification,
   Mycoplasma_result,
   Parameter_code,
   Samplenumber,
   Result
  ) %>%
  dplyr::filter(
   Pathogen_identification %in% c("Mycoplasma bovis")
  dplyr::distinct() %>%
  dplyr::mutate(
   Filenumber_anon = sha256_hash(as.character(Filenumber)),
   Samplenumber_anon = sha256_hash(as.character(Samplenumber))
  dplyr::select(-Filenumber, -Samplenumber)
```

Data manipulation PCR results

```
barometer dt <- barometer dt raw %>%
  dplyr::rename(
   Filenumber=Dossiernummer,
   Samplenumber = Staalnummer,
   Sample_type = Staaltype,
   Parameter_code = PARAMETER_CODE,
   Pathogen = Onderzoek,
   Result = Resultaat,
   Date = Creatiedatum,
   Postal_code = Postcode,
   Farm_ID = ANON_ID
   ) %>%
  dplyr::mutate(
    Country='Belgium',
   Diagnostic_test = case_when(
     Parameter_code %in% c('BAC_AERO','BAC_MYCOPLASMA') ~ 'Culture',
     TRUE ~ 'PCR'
   ),
   Lab reference='1',
   Sample_type = case_when(
     Sample type == "RU Broncho-alveolar lavage (BAL)" ~ 'BAL',
     Sample_type == "RU Anderen" ~'Unknown',
     Sample_type %in% c("RU Swabs", "RU Swab", 'RU Neusswab', 'RU Neusswabs') ~ 'Swab',
     Sample_type %in% c("RU Kadaver", "RU Organen") ~ 'Autopsy',
     TRUE ~ 'Missing'
   ),
   Breed = case_when(
     Bedrijfstype == 'VCALF' ~ 'Veal',
      is.na(MEAT) ~ 'Unknown',
      (as.numeric(MEAT)/as.numeric(TOTAL))>0.9 ~ 'Beef',
      (as.numeric(MILK)/as.numeric(TOTAL))>0.9 ~ 'Dairy',
     TRUE ~ 'Mixed'
   ),
   Pathogen = case_when(
     Pathogen %in% c(
        "AD Pasteurella multocida Ag (PCR)",
        "AD Pasteurella multocida Ag pool (PCR)",
        "AD P. multocida Ag (PCR)",
        "AD P. multocida Ag pool (PCR)") ~ 'Pasteurella multocida',
      Pathogen %in% c(
        "AD Mannheimia haemolytica Ag (PCR)",
        "AD Mannheimia haemolytica Ag pool (PCR)") ~ 'Mannheimia haemolytica',
     Pathogen %in% c("RU PI3 Ag (PCR)", "RU PI3 Ag pool (PCR)") ~ 'PI3',
      Pathogen %in% c("RU BRSV Ag (PCR)", "RU BRSV Ag pool (PCR)") ~ 'BRSV',
     Pathogen %in% c(
        "AD Histophilus somnus (PCR)",
        "AD Histophilus somnus Ag (PCR)",
        "AD Histophilus somnus Ag pool (PCR)",
        "AD Histophilus somni Ag (PCR)",
        "AD Histophilus somni Ag pool (PCR)") ~ 'Histophilus somni',
     Pathogen %in% c(
        "RU Mycoplasma bovis (PCR)",
```

```
"RU Mycoplasma bovis Ag pool (PCR)",
      "RU Mycoplasma bovis Ag (PCR)") ~ 'Mycoplasma bovis',
   Pathogen %in% c("AD Corona Ag (PCR)", "AD Corona Ag pool (PCR)") ~ 'BCV'
 ),
    Province = case_when(
      between(as.numeric(Postal_code), 1000, 1299) ~ 'Brussels',
      between(as.numeric(Postal_code), 1300, 1499) ~ 'Walloon Brabant',
      between(as.numeric(Postal code), 1500, 1999) ~ 'Flemish Brabant',
      between(as.numeric(Postal_code), 3000, 3499) ~ 'Antwerp',
      between(as.numeric(Postal_code), 2000, 2999) ~ 'Limburg',
      between(as.numeric(Postal_code), 3500, 3999) ~ 'Limburg',
      between(as.numeric(Postal_code), 4000, 4999) ~ 'Liège',
      between(as.numeric(Postal_code), 5000, 5999) ~ 'Namur',
      between(as.numeric(Postal_code), 6000, 6599) ~ 'Hainaut',
      between(as.numeric(Postal_code), 7000, 7999) ~ 'Hainaut',
      between(as.numeric(Postal_code), 6600, 6999) ~ 'Luxembourg',
      between(as.numeric(Postal_code), 8000, 8999) ~ 'West Flanders',
      TRUE ~ 'East Flanders'
          )
) %>%
dplyr::select(
 Filenumber,
 Diagnostic_test,
 Samplenumber,
 Country,
 Lab_reference,
 Sample_type,
 Breed,
 Parameter_code,
 Result,
 Pathogen,
 Date,
 Province,
 Farm_ID
) %>%
dplyr::distinct() %>%
dplyr::mutate(
 Filenumber_anon = sha256_hash(as.character(Filenumber)),
 Samplenumber_anon = sha256_hash(as.character(Samplenumber))
dplyr::select(-Filenumber, -Samplenumber)
```

Join all three files

```
barometer <-
barometer_dt %>%
dplyr::left_join(df_samples, by = c('Diagnostic_test','Result', 'Parameter_code')) %>%
dplyr::left_join(
    barometer_aero_cult, by = c('Filenumber_anon', 'Samplenumber_anon', 'Result', 'Parameter_code', 'Samplenumber_anon', 'Result', 'Result
```

```
dplyr::mutate(
 Floored_date = lubridate::floor_date(Date, "month"),
    Pathogen = case_when(
    (Pathogen == 'Pasteurella multocida') ~ 'PM',
    (Pathogen == 'Histophilus somni') ~ 'HS',
    (Pathogen == 'Mannheimia haemolytica') ~ 'MH',
    (Pathogen == 'Mycoplasma bovis') ~ 'MB',
   TRUE ~ Pathogen
 ),
 Pathogen = case_when(
    (Pathogen_identification == 'Pasteurella multocida') ~ 'PM',
    (Pathogen_identification == 'Histophilus somni') ~ 'HS',
    (Pathogen_identification == 'Mannheimia haemolytica') ~ 'MH',
    (Pathogen_identification == 'Mycoplasma bovis') ~ 'MB',
   TRUE ~ Pathogen
 ),
   Result = case_when(
     Result %in% c("Twijfelachtig (PCR)", "POSITIEF", "GEDETECTEERD", "GEDETECTEERD (sterk)", "GEDET.
      "GEDETECTEERD (matig)", "GEDETECTEERD (zeer sterk)", "GEDETECTEERD (zeer zwak)") ~ 1,
     Result %in% c("negatief", "Niet gedetecteerd") ~ 0,
     Result %in% c("NI", "niet interpreteerbaar", "Inhibitie") ~ as.numeric(NA),
    Parameter_code == 'BAC_AERO' & is.na(Pathogen_result) ~ 0,
   Parameter_code == 'BAC_AERO' & !is.na(Pathogen_result) ~ 1,
   Parameter code == 'BAC MYCOPLASMA' & is.na(Mycoplasma result) ~ as.numeric(NA),
   Parameter_code == 'BAC_MYCOPLASMA' & Mycoplasma_result == 'neg' ~ 0,
   Parameter code == 'BAC MYCOPLASMA' & simisc::str contains(Mycoplasma result, 'POS') ~ 1,
   TRUE ~ as.numeric(NA)
) %>%
group_by(
 Lab_reference,
 Country,
 Breed,
 Floored_date,
 Province,
 Farm_ID,
 Diagnostic_test,
 Sample_type,
 Pathogen
 ) %>%
summarise(across(c(Result), max))
```

Save file (long version)

```
write.csv(barometer, "../Data/CleanedData/barometer_DGZ.csv", row.names=TRUE)
```