```
35th INTERNATIONAL CONGRESS OF
MEAT SCIENCE AND TECHNOLOGY 1989
Poster on:
ENZYMATIC BONE CLEANING
AND SCRAP MEAT RECOVERY
N.H. Sørensen & P.B. Rasmussen
Enzyme Process Division
NOVO-NORDISK A/S
DK-2880 Bagsværd - D E N M A R K
INTRODUCTION
As a Supplement or an alton Supplement or an
alternative to traditional
Rendering, enzymatic processes
Rnzyme in developed in Novo's
Maryme Application Pilot Plant
Whereby the following main
products can be made:
  A bland soluble protein
  hydrolysate suitable as
  ingredient in food products
  for human consumption.
0
  A protein hydrolysate
  Suitable for animal feeding.
1
  A completely clean bone
  fraction devoid of meat and
  fat for gelatine production.
SCRAP MEAT RECOVERY
Process 1
Fresh food grade bones are
Crushed to a size of
approximately 20 mm and defatted
by a traditional warm water
extraction method.
The Cleaned crushed bones and defatted wixed
defatted crushed bones and
a lil stickwater are mixed in
<sup>a l:1</sup> ratio and treated with the
<sup>neutral</sup> ratio and treated with the
Neutral Novo protease Neutrase
Paramet Paramet coolied:
parameters are applied:
Enzyme dosage:
                  2-3 kg Neutrase
Temperature
                  0.5 L per ton of
                  wet bones
58-62 C
              •
Reaction time:
                  unadjusted,
                  neutral
                  1 hour
```

n

The hydrolysis is performed in a tank equipped with heat mantle and agitator. The protein hydrolysate is recovered by screening and decanter centrifugation. The superflow from the decanter is fractionated further using a solid injection centrifuge. Should the supernatant contain substantial amounts of fat a fat separation may be necessary before the protein hydrolysate is pasteurized and further concentrated in a falling film evaporator.

In order to produce a completely soluble food grade meat protein powder it is advisable to concentrate to around 25% dry matter, then remove particulate material by filtration before spray drying. A flow sheet on the scrap meat recovery process is presented in Fig. 1.

Products

Besides the fat a low fat/low salt protein powder with a bland non bitter taste is produced. The taste is evaluated from a 20% dry substance solution at pH 6.5. The composition of the protein powder is shown in Table 1. The amino acid composition compared to the one of lean beef and collagen is shown in Table 2. Among the essential amino acids, especially the content of methionine and tryptophane will be low (not measured). But it should be stressed that this product is an ingredient rather than sole nutrient. The content of hydroxyproline is on the other hand high compared to meat. This is due to the fact that collageneous sinew material is hydrolyzed during the enzyme treatment.

Table	1	Typical composition of	
		spray dried food grade	
		protein powder	

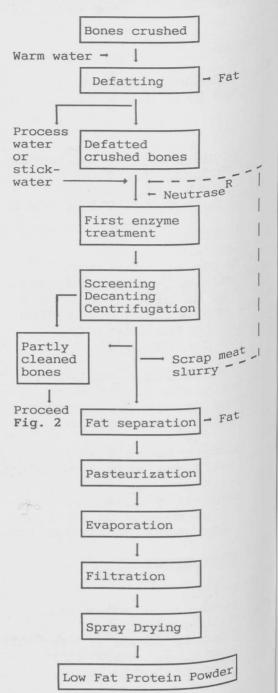
Dry substance (105 ⁰ C)	96%
Protein (%Nx6.25)	80%
Fat (SOXTEC)	18
Ash $(600^{\circ}C)$	88
Average peptide	
chain length (TNBS)	15

Table 2 The composition of essential amino acids and hydroxyproline (g/100 g of protein) in the food grade protein powder compared to lean beef and collagen

	Lean	Protein	Colla-
	Beef	Powder	gen
Isoleucine	4.8	2.5	1.9
Leucine	8.1	5.6	3.5
Lysine	8.9	5.4	4.3
Methionine Phenyl-	2.7	nm*	0.8
alanine	4.4	3.1	2.4
Threonine	4.6	2.8	2.2
Tryptonph.	1.5	nm*	0
Valine Hydroxy-	5.0	3.5	2.7
proline	0	7.4	12.5

* (nm = not measured)

Fig. 1 SCRAP MEAT RECOVERY PROCESS



2

BONE CLEANING

5

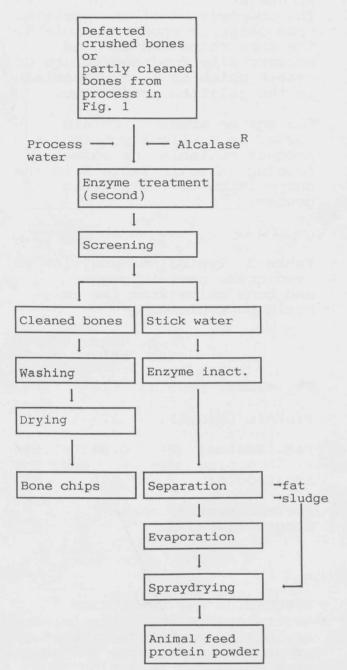
Process 2 This process can be carried out as the second step of the scrap meat recovery process (as indicated in Fig. 1). But if producing food grade protein powder it is recommended that this process takes the place of step one as well. Then instead of having two enzyme treatments there will be only one. It could be the case if raw material without food grade status was to be used in animal feed. The Crushed and defatted bone material is mixed with process water.

Water, e.g. condensate from the evaporator or stickwater, and protease Alcalase 2.4 L. The used.

Enzym

ED Zz
Enzyme dosage : 0.5-1.0 kg
Alcalase 2.4 L
AICAIASE 2.4 L
per ton of wet
bones bones
PH Temperature : 60-70°C
React: unadjusted
tion time : 1 hour
Reaction time : 1 hour This enzymatic treatment secures sinew put release of meat and
d enzimely
a complete release of meat and after protein from the bones and
olnew release of meat and
dit. Protein from the hones and
Separ the reaction the bones are
Watarated from the meat stick
Water by screening.
Way 1-2 cm 1
The 1-2 cm bone fraction is and dried under vacuum to
and dried under vacuum to pprox. 10% water content.
approx. 10% water content.
Ullo
MOT CO the case is a set of the
it heat stall that Alcalase is
in May be stable than Neutrase
it may be necessary to stick wate the enzyme in the
fick wate the enzyme in the
Water i i i e u
reaminutes. After separation
inactivate stable than Neutrase inactivate the enzyme in the for 6 water by heating to 85°C of residual sludge and fat the spraydried. A flow sheet on the
^{spraydried} . A flow sheet on the ^{presented} process is
bondydried is evaporated and
Pro Cleani A ILOW Sheet on the
esented process is
bone cleaning process is presented in Fig. 2.

Fig. 2 BONE CLEANING PROCESS



Products

The composition of the purified bone chips is shown in Table 3. The bone chips can be used commercially for manufacture of ossein which is a raw material in the gelatine production.

The enzyme treated protein powder is a slightly bitter product suitable for animal feeding. Also in Table 3 is the composition of the protein powder.

Table 3 Typical composition of feed grade protein powder and bone chips from the enzymatic bone cleaning

		Protein powder
Dry subst. (105 ⁰ C)	91%	96%
Protein (Nx6.25)	27%	84%
Fat (Soxtec)	0.8%	1.6%
Ash (600 ⁰ C)	59%	98
Average peptide cha Length (TNBS)	6	

DISCUSSION AND CONCLUSION As it appears from this paper we are having an alternative of two processes, a two step and an one step enzymatic rendering process.

The chosen process depends on

the quality of the raw material and the quality of protein which is in demand.

The two step process is only profitable when the protein can be sold as a food grade product meant for human consumption.

Technically the processes have been designed in a way so that they can easily be carried ^{OUT} in a normal slaughterhouse or rendering plant.

Considering the enzyme Neutrase the chosen temperature, pH, dosage and reaction time ensure a high initial activity of the enzyme and that after one hour the enzyme has been completely inactivated. In that way the degree of hydrolysis is limited to around 5%, ensuring a nonbitter protein hydrolysate.

Conclusively, NEUTRASE 0.6 L ¹⁵ recommended for use in a twostep scrap meat recovery process where the products are primarily food grade proteins for human consumption and bone chips gelatine production.

Where ALCALASE 2.4 L is used the protein product is most suitable as an animal feed ingredient.

These processes have been developed with the aim of demonstrating a few of the man^y possibilities which exist when biotechnology is introduced food technology. Thereby chalenging the industry to obtain inspiration for further developments.

Fig. 1 SCRAP MEAT RECOVERY PROCESS

1 Ch

n t

3

3

į

3

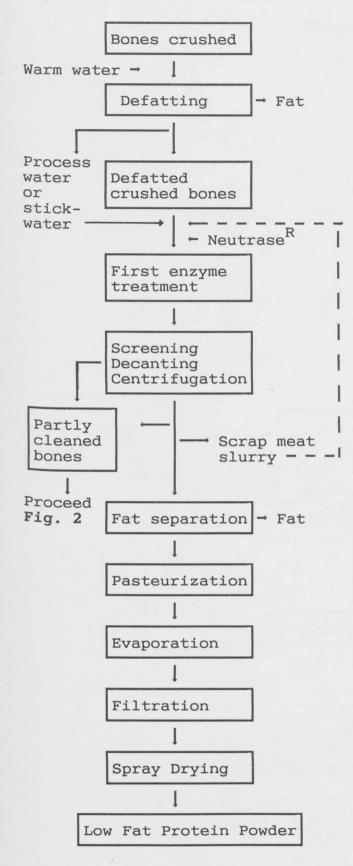
35 L¥

10

e

1

0



961

Fig. 2 BONE CLEANING PROCESS

