

Antibodies Collection dedicated to food components

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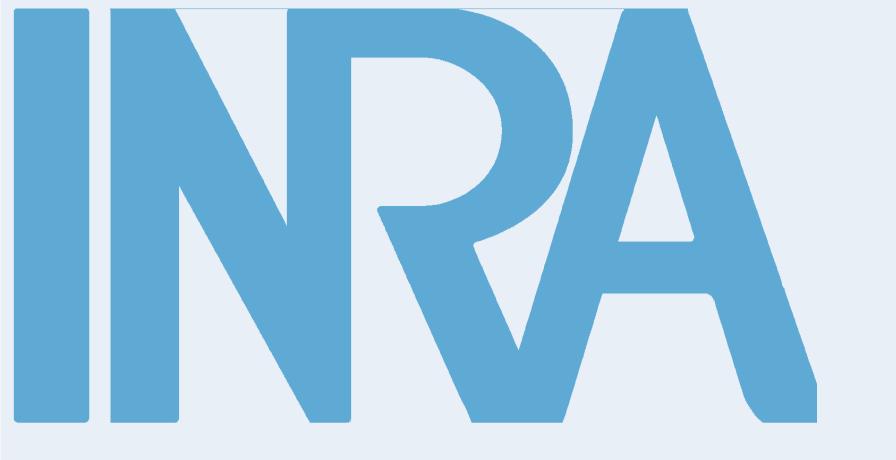
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Antibodies Collection dedicated to food components

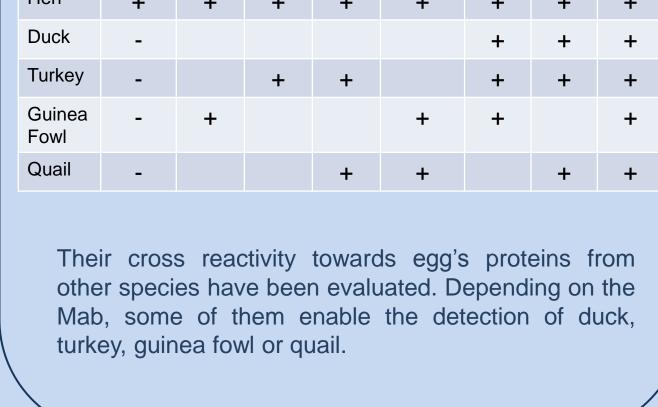
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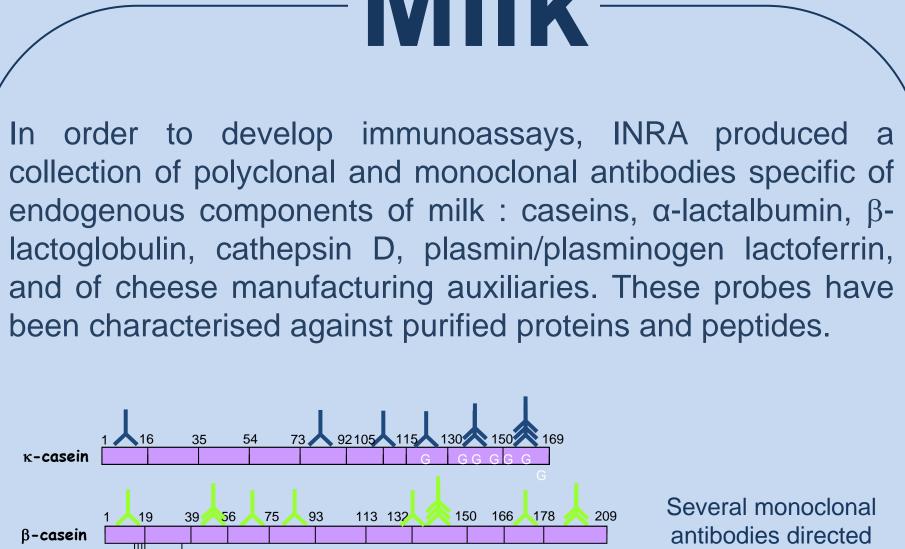
¹UR 1268 Biopolymères Interactions Assemblages INRA Nantes, France; ²UR 342 Technologie et Analyses Laitières INRA Poligny, France; ³UR 1253 Science et Technologie du Lait et de l'Oeuf INRA Agrocampus Ouest Rennes, France

Due to their high selectivity, antibodies are useful tools to detect, quantify or characterise proteins or peptides from complex samples. Numerous antibodies against wheat, milk and egg components were obtained at INRA. This large library of antibodies enables the diversity of composition of raw material and impact of processes and digestion to be addressed.

Cereals Antibodies against wheat grain proteins i.e. proteins from gliadin, glutenin and albumin/globulin fractions have been produced from purified proteins or synthetic peptides. Antibodies have been fully characterised by ELISA with purified prolamins proteins Some of these antibodies enable ONT 18A51D2 gliadins or glutenins subclasses to be distinguished from one another, others are suitable for S1 EG7D11 the detection of the whole gliadin S1 DH3 or glutenin fractions or whole S1 EG87 prolamins. Combining Macro-Array spotting chemiluminescent ELISA, the Wheat Antibody Collection was screened against « prolaminlike » proteins from several cereals. Most of the antibodies bound proteins from Interestingly some of them also reacted on brachypodium, the new model plant for **Dilution of Antibody PQQ 3B4**

Numerous monoclonal antibodies against purified proteins from hen's egg have been produced. Antibodies specific of ovalbumin, ovomucoid, ovotransferrin and lysozyme from the white and phosvitin, LDL and HDL from the yolk have been fully characterised and are now available for characterisation studies of raw material. | Grp 1 | Grp 2 | Grp 3 | Grp 4 | Grp 5 | Grp6 | Grp7 | Grp 8 | I Mab | I Mab

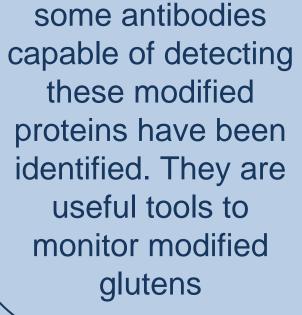


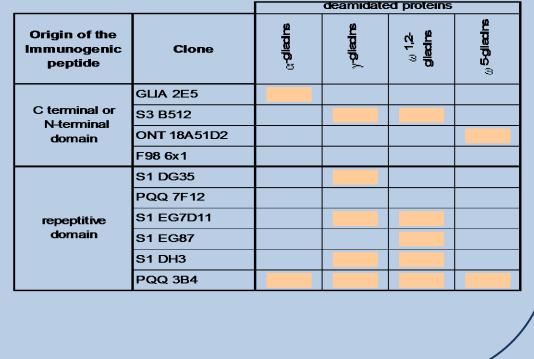


Several monoclonal antibodies directed against β , κ , α_{s1} and α_{s2} caseins were produced. Specificities were determined using libraries of synthetic peptides and peptides fractionated from tryptic hydrolysis of purified caseins.

Food processes may have an important impact on protein structure and induce chemical changes. Biological activities and technological properties of these processed products can then be affected as well as the detection of their components by antibodies. Within the collection some antibodies enable to detect these modified proteins.

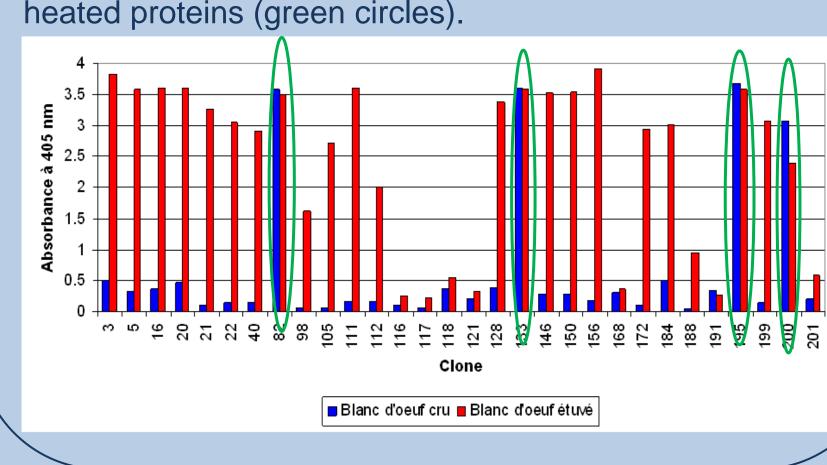
Wheat deamidation In some food products, gluten is added after partial hydrolysis or deamidation in order to enhance their use as functional ingredients. These processes lead to the replacement of some glutamine by glutamic acid which may impaire the immunochemical detection of the wheat prolamins.





Egg heating

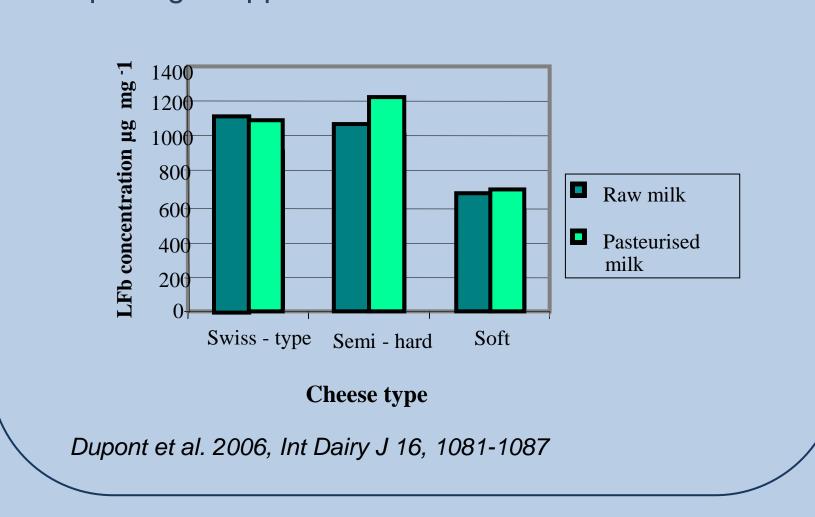
Heating can affect structure of proteins, lead to partial hydrolysis or induce chemical modifications. Antibodies can be sensitive to these changes. Among the numerous antibodies directed against egg's proteins, some of them were selected for their abilities to detect both native and heated proteins (green circles).



Cheese ripening

After selection of the appropriate antibodies a sandwich ELISA was developed for lactoferrin quantification in milk and cheese.

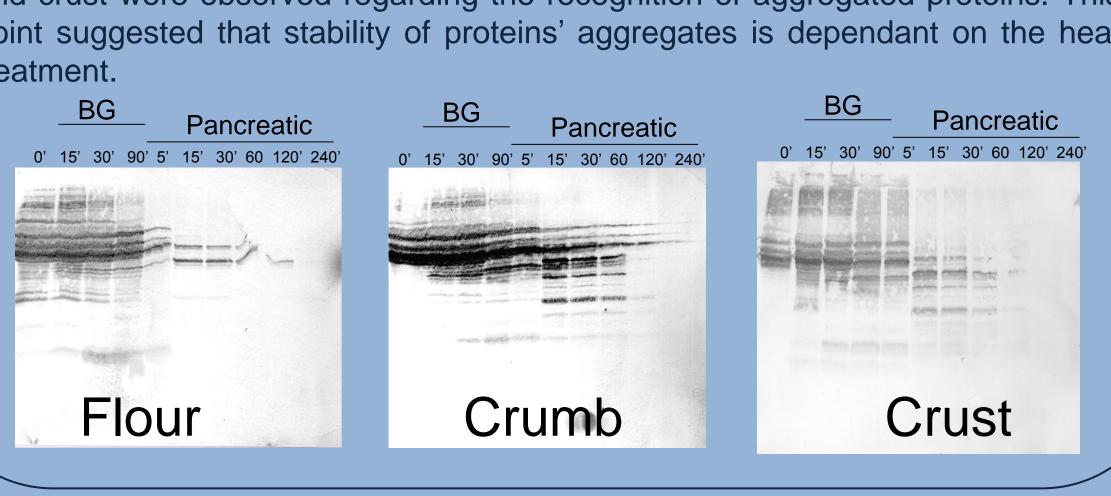
Lactoferrin is present in all cheese types at the end of ripening in appreciable amounts.



Antibodies enable the monitoring of the degradation of particular structure during digestion. Antibodies to wheat and milk were successfully used to evaluate the impact of some food process on the in vitro digestion of some peptides implied in food allergies to wheat and to milk.

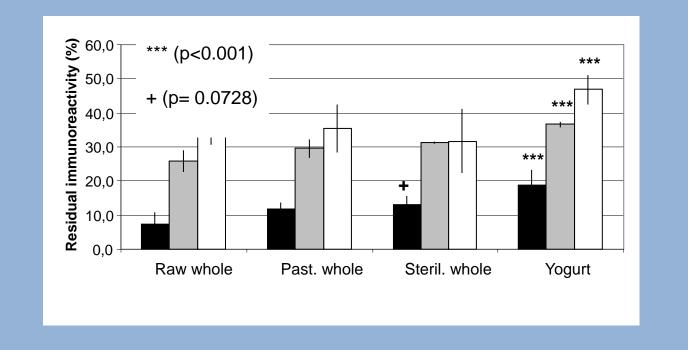
Wheat digestion eating on in vitro Bucco-Gastric (BG) and pancreati

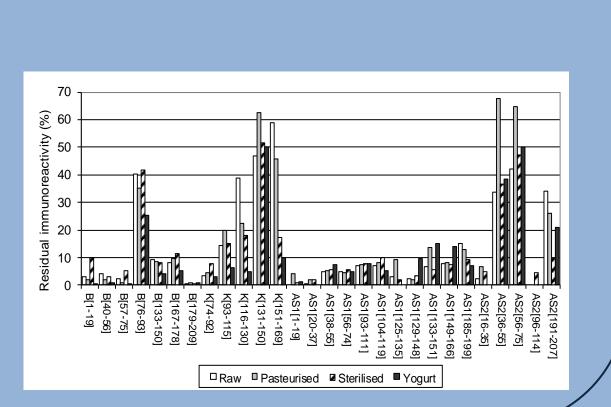
Influence of heating on *in vitro* Bucco-Gastric (BG) and pancreatic digestion of wheat flour, bread crump and crust was evaluated with antibodies specific to C-terminal peptide of β -gliadin by western blotting. Heating slow down the digestion but did not seem to change the immunoreactivity. Differences between crumb and crust were observed regarding the recognition of aggregated proteins. This point suggested that stability of proteins' aggregates is dependant on the heat treatment.



Milk digestion

Different dairy matrices were manufactured and subjected to *in vitro* digestion using an infant gut model. Among other technics, hydrolysis of caseins (CN) was evaluated with immunochemical techniques. Residual immunoreactivity assessed by inhibition ELISA revealed that heat treatment increases the resistance of CN to infant digestion. Thanks to the set of monoclonal antibodies covering the sequence of caseins, parts of the different caseins which were resistant to digestion were identified.





Precise characterisation of the antibodies and their gathering in a unique collection make this antibodies' library a genuine toolbox to address raw material diversity, molecular complexity and the impact of process and digestion on food component and architecture.