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**Teams at IFPEN studied the phenomenon of fuel autoxidation, phase separation and deposit formation causing a variety of malfunctions. Having proposed a new deposit formation process, researchers validated it for diesel, biodiesel and aviation kerosene fuels using a new methodology and demonstrated the role of charge carriers and polarity at the oxidized/non-oxidized product interface.**

### **Autoxidation, a source of deposits and malfunctions**

Autoxidation refers to the phenomenon whereby a fuel tends **to break down as a result of the oxidation of its compounds at low temperature** (generally  $T < 300^{\circ}\text{C}$ ). This form of fuel degradation may result in **the formation of insoluble products and deposits of diverse origin and compositions in tanks, combustion**

systems and injection circuits.

These deposits can then lead to **mechanical malfunctions** affecting pumps and injectors, filter clogging or **malfunctions relating to increased pressure losses** through filters.

Consequently, fuel stability in the context of autoxidation is a major issue. It concerns the entire logistics chain from storage through to its transportation and use in different modes of land, air and naval transport.

### How are these deposits formed?

IFPEN has built up **solid expertise in the field of fuel stability** by studying deposit formation mechanisms. The objective: to gain a better understanding of **the liquid phase autoxidation process** and develop detailed, semi-detailed and global kinetic models designed to predict both fuel degradation as a function of its chemical composition and operating conditions as well as deposit formation.

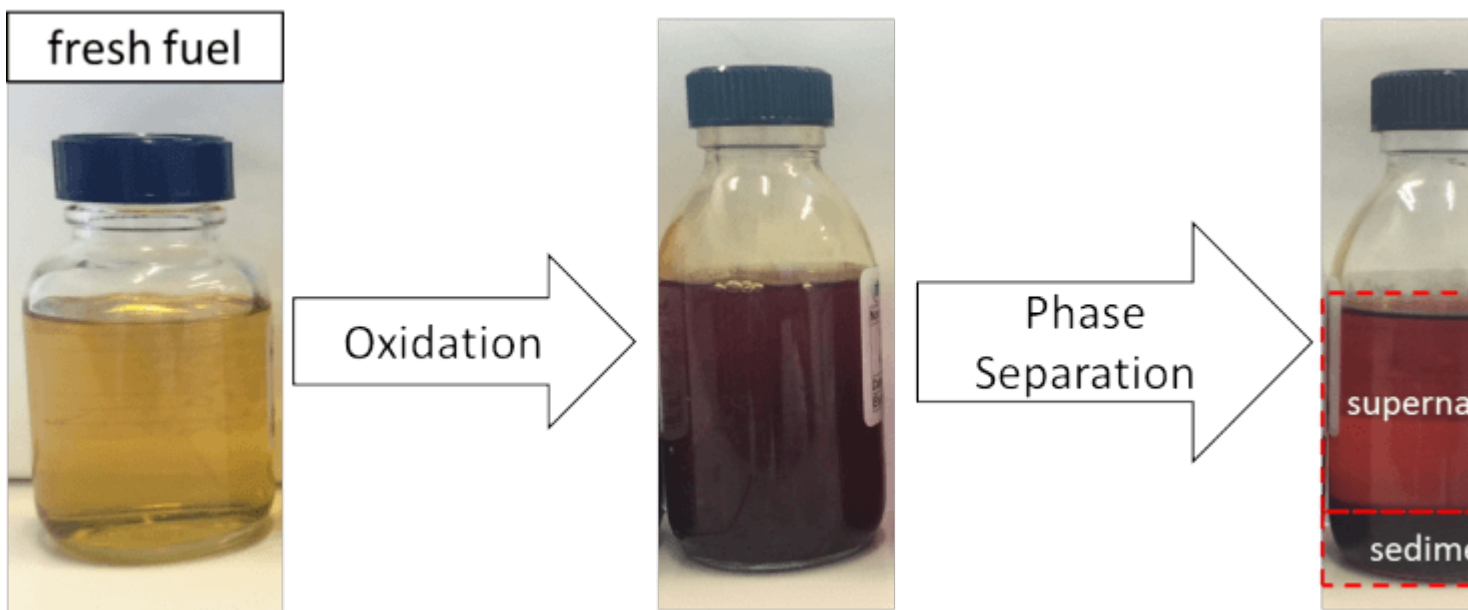


Figure 1. Photo of a biodiesel before and after oxidation and monitoring of phase separation with the formation of a supernatant and a sedimented phase [1]

Research conducted in recent years at IFPEN has mobilized a broad range of internal expertise<sup>1</sup> and made it possible to demonstrate that deposits are primarily formed **during the engine and turbine stop phases**[2].

Accelerated oxidation methods employed in the laboratory<sup>2</sup> involving real and model (surrogates) fuels made it possible to observe **a liquid-liquid separation phase as a function of fuel oxidation conditions** (figure 1). Detailed analyses of both the supernatant fractions and sedimented phases, particularly via GC-MS<sup>3</sup>, led to the identification of **the broad chemical families of stable products** formed during autoxidation processes: ketones, epoxides, aldehydes, alcohols and carboxylic acids[3–5]. These products appear to play an important role **in the formation of several deposit precursors**.

**The characteristics of the aggregates do not, in themselves, explain phase separation**

Research teams have exploited the complementary nature of molecular characterization techniques, such as GC-MS and FTIR<sup>4</sup>, and other approaches, such as **low field NMR and X-ray scattering** in order to improve our understanding of such mechanisms [6].

By way of example, small-angle scattering techniques, SAXS<sup>5</sup> and SANS<sup>6</sup>, made it possible **to characterize the aggregation mechanisms demonstrated** - diffusion limited aggregation (DLA) and reaction limited aggregation (RLA)[5] - as well as **to monitor aggregate size**, which increases in a linear manner with the number of particles in test conditions. Nevertheless, the physical characteristics alone (particle size and number) are not thought to explain the observed phase separation and deposits.

### **Polarity differences between oxidation products and the non-oxidized fluid**

In order to identify the principal levers leading to phase demixing, IFPEN's teams used **optical microscopy** as well as **Turbiscan methods** and FTIR[1]. Figure 2 shows the evolution via demixing of oxidized samples of a complex emulsion. Turbiscan results revealed that the demixing rate was not directly linked to aggregate density or viscosity. Instead, the phenomenon seemed to be primarily related to **differences in polarity** (due for example to oxygenated compounds) of the sedimentary and supernatant phase content, which was confirmed by the FTIR results.

# Oil-oil system after fuel oxidation: before phase separation

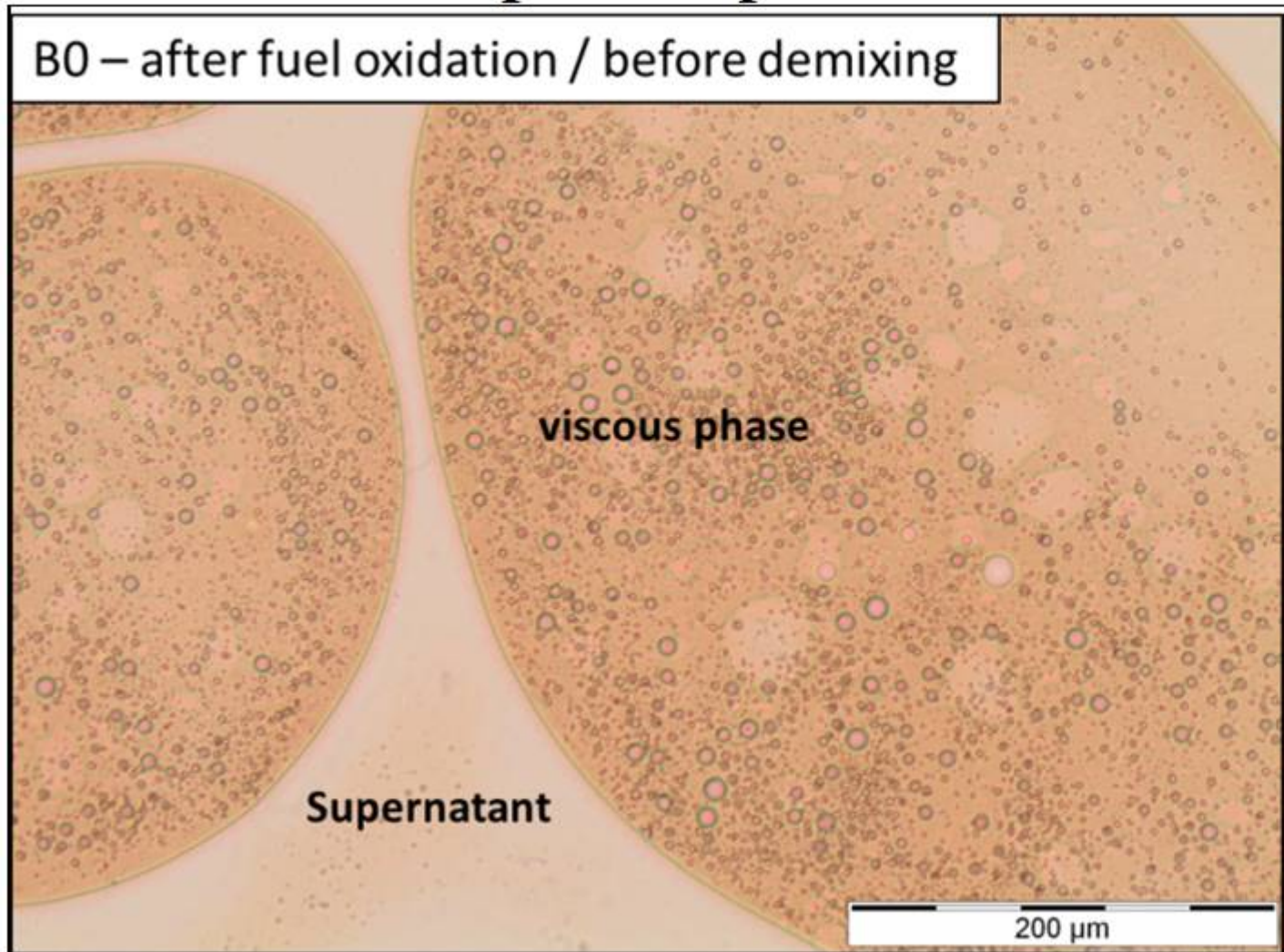


Figure 2. Optical microscope images of an oxidized diesel prior to phase separation (left) and the sedimented phase after phase separation (right) [1]

Everything seems to indicate **that deposit precursors are formed in the liquid phase**, and, therefore, that the deposit is not directly generated by the surface in contact with the fuel. These observations led to the proposal of **a new deposit formation mechanism** [1] whereby the difference in polarity between the oxidation products and the non-oxidized fluid represents a significant phase separation lever (figure 3).

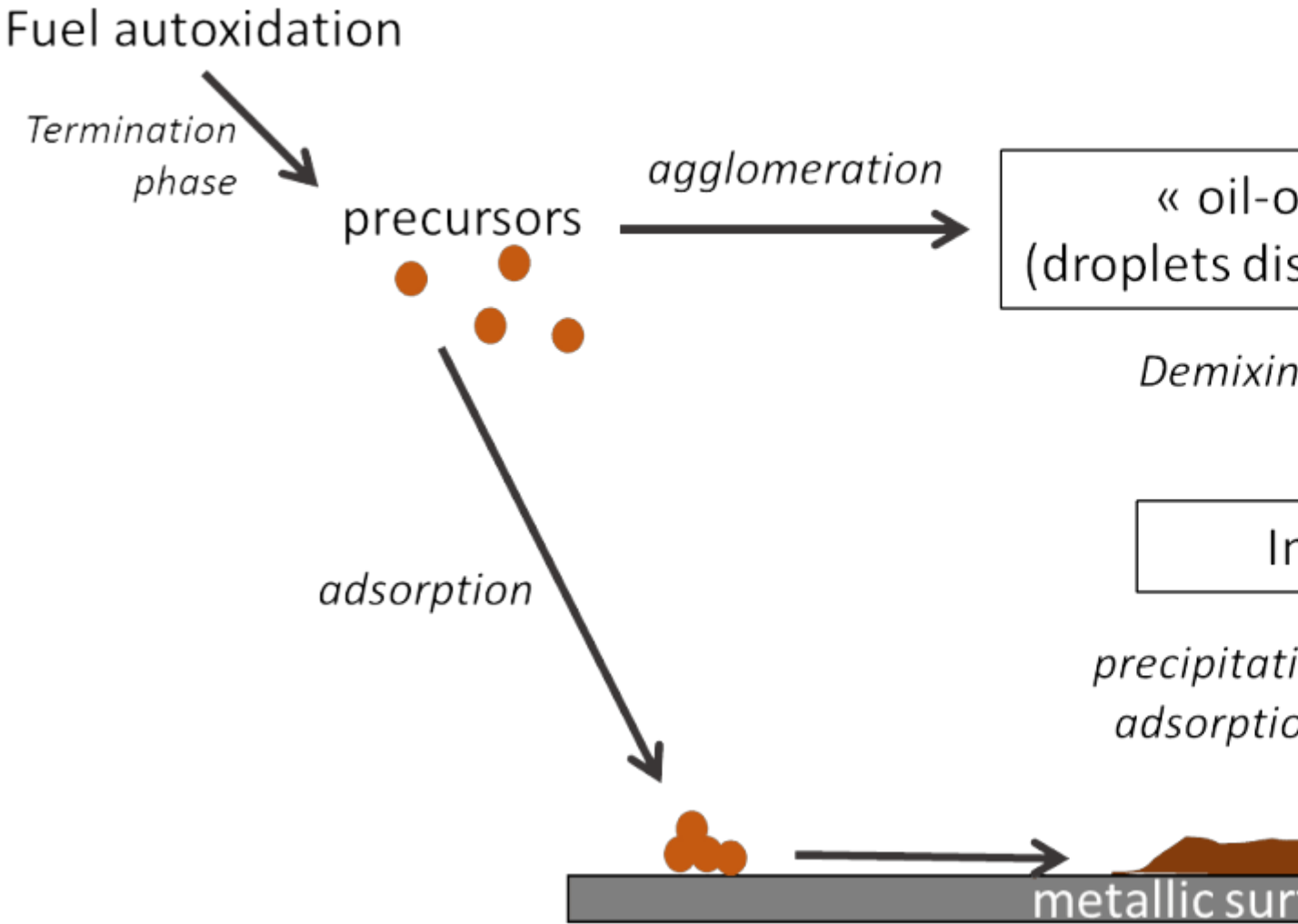


Figure 3. Proposed mechanism process for the formation of deposits resulting from fuel degradation [1]

### IFPEN validates a new deposit formation mechanism

To validate this mechanism proposal, experimental research was conducted using **electrochemical impedance spectroscopy**, combined with an **electrochemical quartz crystal microbalance (EIS/EQCM)**. These combined techniques make it possible to analyze the dynamic behavior of a system. The idea is to apply an electrical disturbance to the system over time and monitor its response, as well as the mass of the deposit formed. This methodology highlighted **the role of charge carriers and polarity at the oxidized/non-oxidized product interface** on the deposit formation process for diesel, biodiesel and aviation kerosene fuels. Moreover, due to its efficacy, **this methodological application has resulted in the filing of a patent**[7].

In a part of the study dedicated to aviation fuels, EIS/EQCM coupling highlighted different processes leading to deposit formation on the microbalance from sulfur and aromatic compounds present in Jet A1. **Capacitance** (i.e., the polarity at the oxidized/non-oxidized product interface) seems to be the main parameter determining deposit formation.

This research has helped establish IFPEN as one of the leading players in the field of fuel stability in France and internationally.

<sup>1</sup> Chemical analysis, structural analysis and imaging, complex fluids, colloids and condensed matter, surface, interface and materials science

<sup>2</sup> PetroOxy or RSSOT, Rancimat, and Parr Autoclave

<sup>3</sup> Gas chromatography (GC)-Mass spectrometry (MS) coupling

<sup>4</sup> Fourier-Transform Infrared Spectroscopy

<sup>5</sup> Small-Angle X-Ray Scattering

<sup>6</sup> Small Angle Neutron Scattering

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