Press release

Availability of a first document in response to questions received from shareholders

LYON, France, October 20, 2025

POXEL SA (Euronext: POXEL - FR0012432516), informs that a first document in response to the questions received from the shareholders is available in French on the Company's website, in the **"Investors" section, "Shareholder Info", "Documentation"**.

This first document covers the main themes raised by shareholders over the past few months, concerning:

- the strategy of partnerships established in the past,
- the current relationships of Poxel with Merck Serono,
- the business partnership with Sumitomo for TWYMEEG® in Japan, and
- the rights related to Roivant.

The publication of this document is part of the new governance's desire to strengthen communication with all its shareholders and establish a constructive dialogue with them.

About Poxel

Poxel is a **clinical stage biopharmaceutical company** developing **innovative treatments for chronic serious diseases with metabolic pathophysiology**, including **metabolic dysfunction-associated steatohepatitis (MASH)** and rare disorders. For the treatment of MASH, **PXL065** (deuterium-stabilized *R*-pioglitazone) met its primary endpoint in a streamlined Phase 2 trial (DESTINY-1). In rare diseases, development of **PXL770**, a first-in-class direct adenosine monophosphate-activated protein kinase (AMPK) activator, is focused on the treatment of adrenoleukodystrophy (ALD) and autosomal dominant polycystic kidney disease (ADPKD). **TWYMEEG®** (Imeglimin), Poxel's first-in-class product that targets mitochondrial dysfunction, is now marketed for the treatment of type 2 diabetes in Japan by Sumitomo Pharma and Poxel expects to receive royalties and sales-based payments. Poxel has a strategic partnership with Sumitomo Pharma for Imeglimin in Japan. Listed on Euronext Paris, Poxel is headquartered in Lyon, France, and has subsidiaries in Boston, MA, and Tokyo, Japan.

For more information, please visit: www.poxelpharma.com

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