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Poxel: Availability of a Second Document Addressing Questions Received From Shareholders

LYON, France--(BUSINESS WIRE)-- Regulatory News:

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

This new document summarizes four other topics raised by shareholders in recent months:

- Trusts
- Financing
- Debts and loans, and
- Stock market and shareholders.

The publication of this document reflects the new governance's commitment to enhancing transparency and fostering an open, constructive dialogue with all shareholders.

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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