

DATE:	17 June 2024
TO:	All Zones – Physicians, Nurses, Pharmacists, and Managers
FROM:	Clinical Biochemistry, Alberta Precision Laboratories (APL)
RE:	Realignment of <i>Infliximab Levels</i> with the World Health Organization Standard

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Key Message

- Effective **June 17, 2024**, the Immundiagnostik (IDK) *Infliximab Level* enzyme linked immunosorbent assay (ELISA) calibration will realign with the National Institute for Biological Standards and Controls (NIBSC) *Infliximab* World Health Organization (WHO) standard.

Background

- Over the past year, Alberta Precision Laboratories has been working closely with Affinity Diagnostics to correct calibration drift especially for *Infliximab* levels above the therapeutic range.

How this will impact you

- The new lot of calibrators are expected to produce a negative proportional bias (i.e., on average shift *Infliximab* levels down by 25%). The shift will be more noticeable at *Infliximab* levels above 10ug/mL.
- An *Infliximab Level* comment will append in the report to alert physicians.

Action Required

- Known inflammatory bowel disease patients undergoing *Infliximab Level* therapeutic drug monitoring will require re-baselining.

Questions/Concerns

- Dr. Kareena Schnabl, Clinical Biochemist, UAH Special Chemistry Lab: 780 407-3186;
Kareena.schnabl@aplabs.ca

Approved by

- Dr. Allison Venner, Section Chief Clinical Biochemistry, South Sector, APL
- Dr. Michael Mengel, Medical Director, North Sector, APL
- Dr. Dylan Pillai, Medical Director, South Sector, APL

Effective September 1, 2023, APL has become the sole provider of all public lab services in Alberta. As a result, community lab services formally provided by DynaLIFE Medical Labs will become the responsibility of Alberta Precision Labs (APL). This change impacts all zones.