

## Factsheet

# Epstein-Barr Virus–Associated Post-Transplant Lymphoproliferative Disease or Disorder (EBV<sup>+</sup> PTLD)

*A rare, life-threatening complication following transplant*

## Overview

EBV<sup>+</sup> PTLD is a serious condition that can develop in patients who have received a **solid organ transplant** (such as a kidney, liver, heart, or lung) or an **allogeneic stem cell (bone marrow) transplant**.

To prevent rejection, transplant recipients must take medications that suppress their immune system. This immune suppression can make patients susceptible to the **Epstein-Barr virus (EBV)**, a common virus most people carry. In some patients, this virus may grow unchecked and can lead to a cancer known as post-transplant lymphoproliferative disease or disorder (PTLD).

## The role of Epstein-Barr virus

- EBV is one of the most common human viruses
- In people with healthy immune systems, EBV is kept under control
- After transplant, suppression of **T cells** weakens this control
- Loss of immune surveillance is a key driver of EBV<sup>+</sup> PTLD

## Who is affected

EBV<sup>+</sup> PTLD can affect both:

- Solid organ transplant recipients
- Stem cell (bone marrow) transplant recipients

Key risk considerations:

- PTLD occurs in approximately **2–20%** of transplant recipients, depending on transplant type
- Risk is highest:
  - » Within the **first year after transplant**
  - » Again **5–15 years after transplant**
- Risk is generally higher in solid organ transplant recipients

## Severity and outcomes

EBV<sup>+</sup> PTLD is an aggressive disease with poor outcomes, particularly when it does not respond to treatment or returns after initial therapy.

- Mortality rates:
  - **70–90%** following stem cell transplant
  - **50–70%** following solid organ transplant
- Median survival:
  - **Less than one month** for stem cell transplant recipients
  - **Approximately four months** for solid organ transplant recipients

## Current treatment landscape

There are **no FDA-approved therapies** specifically for EBV<sup>+</sup> PTLD.

Current approaches may include:

- Reducing immune-suppressing medications, which may increase the risk of organ rejection
- Antibody-based therapy, sometimes combined with chemotherapy

Limitations of current care:

- Variable treatment landscape
- High risk of infections and treatment-related toxicities
- Many transplant patients cannot tolerate aggressive therapy due to immune suppression

## Unmet medical need

Patients with EBV<sup>+</sup> PTLD face:

- Few treatment options
- High mortality rates
- Very poor outcomes when disease is relapsed or refractory

Despite the severity of the disease, **no therapies are currently approved by the FDA** specifically for EBV<sup>+</sup> PTLD. New, targeted therapies are urgently needed.

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