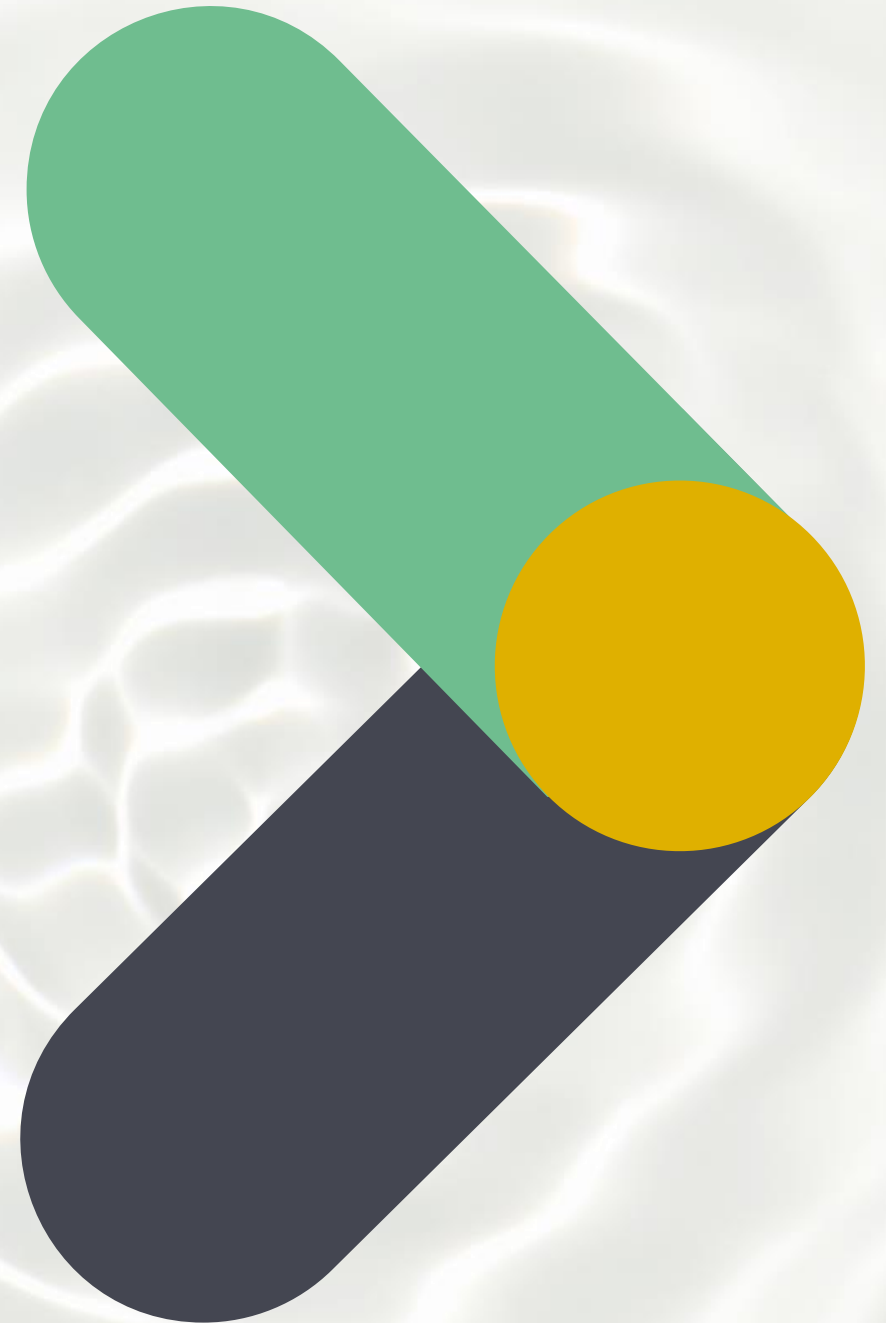




# NATEVBA

(vevasumab)

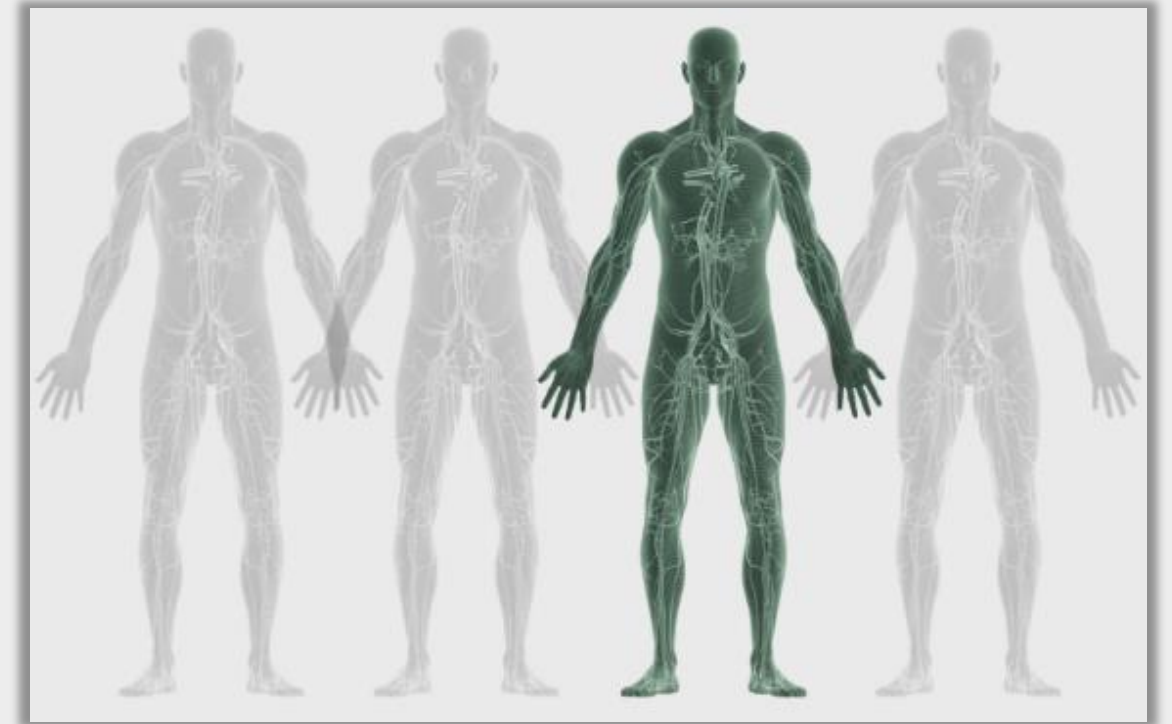
Natevba (vevasumab):  
Precision targeting in XA+  
Non-Hodgkin's lymphoma



Current treatments do not effectively target malignant cells and therefore have high toxicity<sup>1</sup>

Unlike in other types of NHL, monoclonal antibody treatments show poor efficacy in XA+ subtypes and are generally not used<sup>1</sup>

Chemotherapy is the standard of care, but has limited efficacy and high toxicity<sup>1</sup>



**1 in 4** XA+ NHL patients discontinue chemotherapy due to adverse events.<sup>1</sup>

NHL = non-Hodgkin's lymphoma; XA + = X-antigen-positive.

1. Mcguire T et al. Cancer Therapy Review 2018;12 (4):469–470.

## Natevba offers clinically meaningful improvements in QoL<sup>1</sup>

Natevba improved or maintained QoL for the duration of treatment in all five EuroQoL domains:



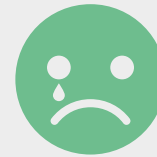
Mobility



Self-care



Usual activity



Pain



Depression

1. Vincent A et al. Cancer Therapy and Management 2020;22 (1):87–88.

QoL = quality of life

# NATEVBA

(vevasumab)

Natevba is an innovative new treatment targeting X-antigen<sup>1</sup>

Natevba is indicated for the first-line treatment of X-antigen positive (XA+) non-Hodgkin's lymphoma<sup>1</sup>



## With Natevba, precision targeting ensures minimal toxicity



Natevba has a more favourable tolerability profile than chemotherapy, with a lower incidence of Grade 3–4 adverse events<sup>1,2</sup>



Natevba has a low incidence of haematological adverse events, including neutropenia<sup>1</sup>



Adverse events are generally reversible with appropriate dose adjustments<sup>1</sup>

1. Vincent A et al. Cancer Therapy and Management 2020;22 (1):87–88.

2. Natevba (vevasumab) Summary of Product Characteristics.

# With Natevba, precision targeting delivers exceptional results<sup>1</sup>

Over 2 years median  
progression-free survival<sup>1</sup>

Meaningful improvements  
in quality of life<sup>1</sup>

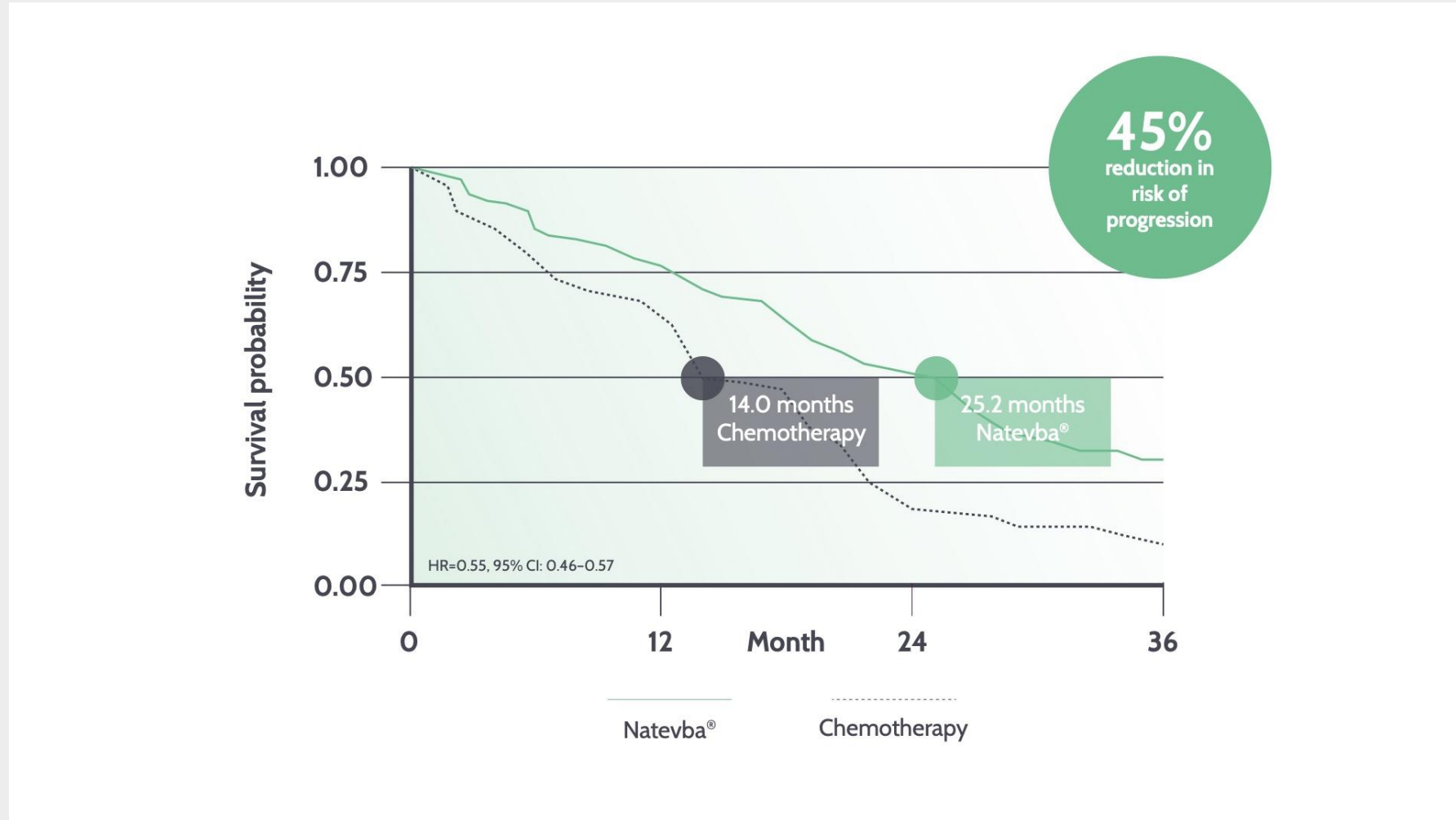


Over 80% response rate<sup>1</sup>

PFS= Progression-free survival QoL = Quality of Life

1. Vincent A et al. Cancer Therapy and Management 2020;22 (1):87–88.

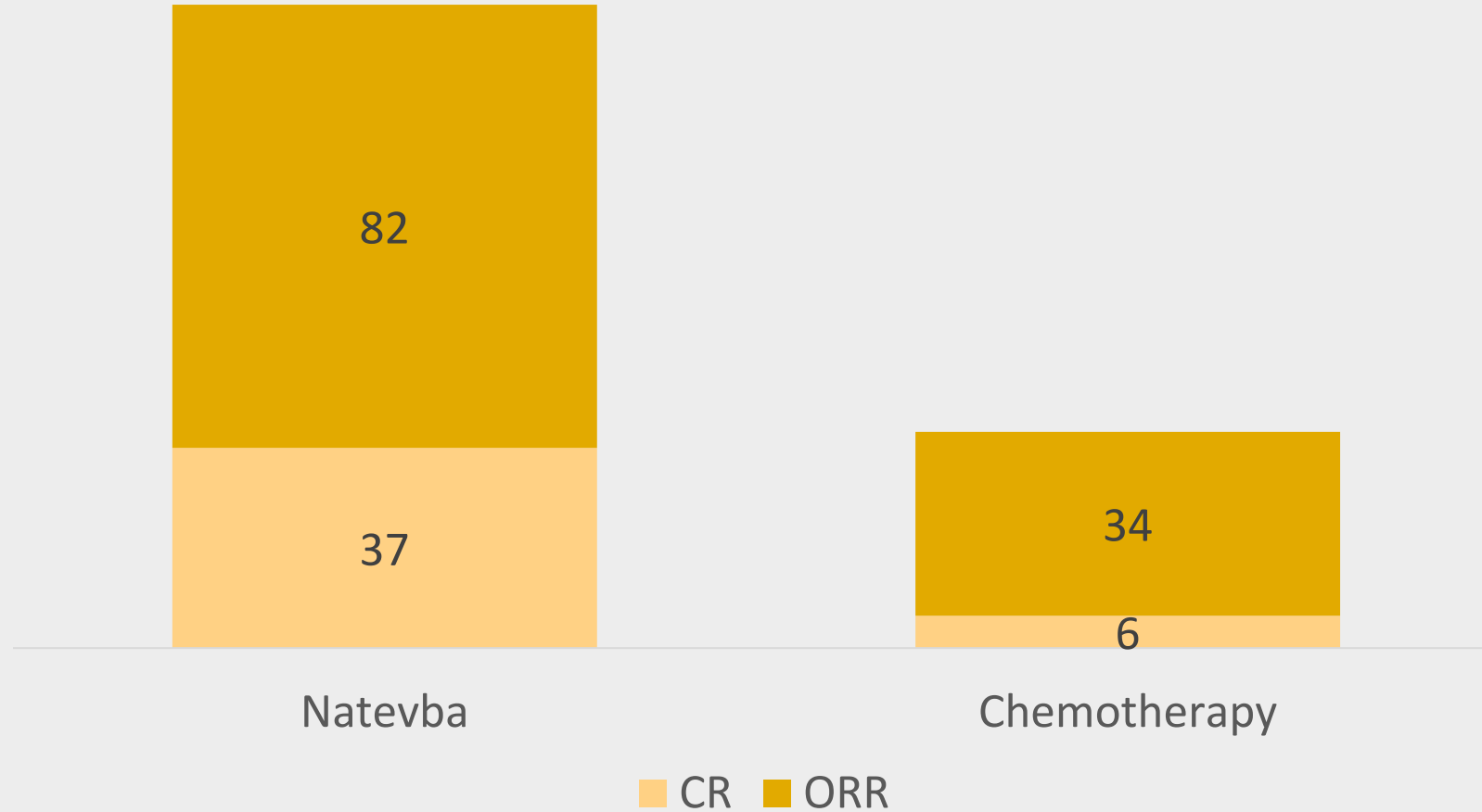
# Natevba delivers over two years median PFS<sup>1</sup>



1. Vincent A et al. Cancer Therapy and Management 2020;22 (1):87-88.

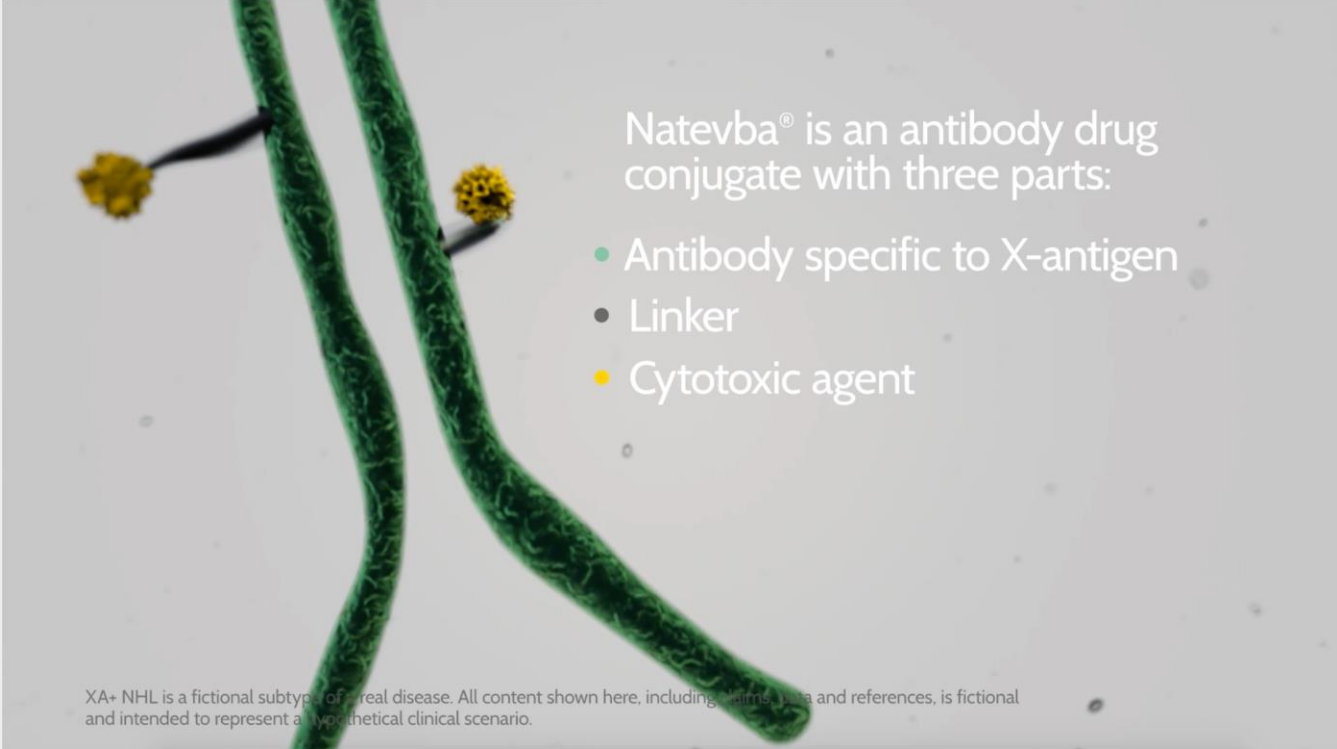
CI = confidence interval HR = hazard ratio PFS = progression-free survival

## Over 80% of patients respond to Natevba<sup>1</sup>



1. Vincent A et al. Cancer Therapy and Management 2020;22 (1):87–88.  
CI = confidence interval CR = complete response ORR = objective response rate

# Natevba is the only treatment to target X-antigen, a highly specific marker of malignant lymphocytes<sup>1</sup>

The diagram shows two green, textured, rod-like structures representing the antibody component of the Natevba conjugate. Each rod has a small black linker attached to its side, which in turn connects to a yellow, star-shaped cytotoxic agent. The background is a light gray with some faint, out-of-focus spots.

Natevba® is an antibody drug conjugate with three parts:

- Antibody specific to X-antigen
- Linker
- Cytotoxic agent

In X-antigen-positive (XA+) non-Hodgkin's lymphoma (NHL), malignant lymphocytes are identifiable by a unique surface marker known as X-antigen.

Natevba is an antibody-drug conjugate with three parts:

- Antibody specific to X-antigen
- Linker
- Cytotoxic agent

Natevba recognizes X-antigen on malignant lymphocytes and binds strongly. Binding triggers uptake of the drug into the cell. Inside the cell, the linker is cleaved, releasing the cytotoxic agent. This induces apoptotic cell death.

XA+ NHL is a fictional subtype of a real disease. All content shown here, including claims, data and references, is fictional and intended to represent a hypothetical clinical scenario.

# Natevba's manageable dosing schedule lets patients get on with life<sup>1</sup>



Natevba has a convenient 4-weekly infusion schedule<sup>1</sup>



The recommended dose is 1 .8 mg/ kg administered as an intravenous infusion over 30 minutes every 4 weeks

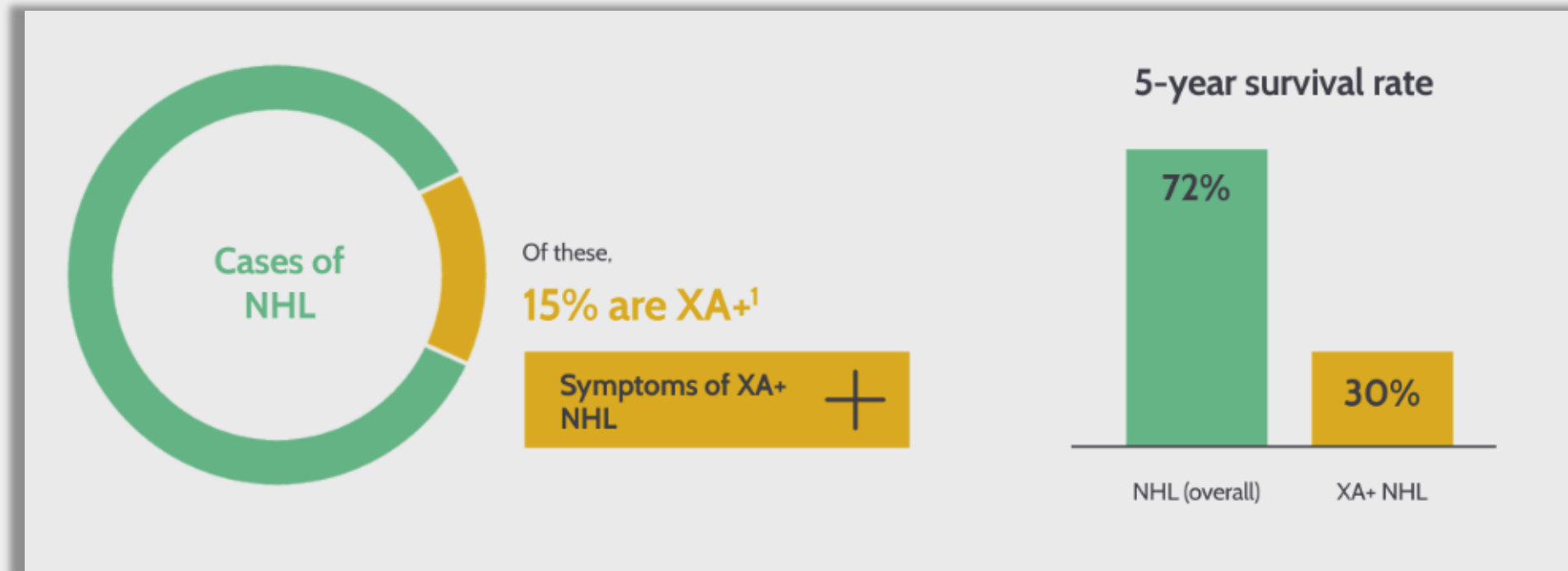


Natevba is indicated as monotherapy, freeing patients from the burden of chemotherapy<sup>1</sup>

# X-antigen positive (XA+) non-Hodgkin's lymphoma (NHL) is a particularly aggressive subtype of NHL, with a poor prognosis

Worldwide, over half a million people are diagnosed with NHL per year<sup>1</sup>

XA+ NHL is an aggressive subtype with rapid progression and a low survival rate<sup>1</sup>



1. Mcguire T et al. Cancer Therapy Review 2018;12 (4):469–470.

## Natevba:

### Precision targeting in XA+ NHL



An innovative new treatment  
targeting X-antigen<sup>1</sup>



Exceptional clinically proven results<sup>2</sup>



Minimal toxicity compared to chemo<sup>2</sup>



NHL = non-Hodgkin's lymphoma; XA+ = X-antigen-positive.

1. Natevba (vevasumab) Summary of Product Characteristics.

2. Vincent A et al. Cancer Therapy and Management 2020;22 (1):87–88.

# Natevba Efficacy

## *Local Event*

Location: Elite Spire Hotel

Date: 15<sup>th</sup> of March 2024

Do you want to attend?

Register | [here](#) |

 **NATEVBA**  
(vevasumab)

