

### Exhibitors' Contact Information

#### **Alexion Contacts:**

> Lisa Saxe

Lisa.Saxe@alexion.com

#### **Amgen Contacts:**

> Jason Bronson

jbronson@amgen.com 405-740-2287

> Jennifer Mason

jmason@amgen.com 918-408-7227

#### **AstraZeneca Contacts**

> Jason Ophus

Jason.ophus@astrazeneca.com 918-693-0662

#### **Aurinia Contacts**

> Chris Reynolds

Immunology Specialist creynolds@auriniapharma.com 405-202-9192

> Catherine Thorson

Regional Professional Relations cthorson@auriniapharma.com 405-833-1641

> Sam Smothers

Field Access Navigator ssmothers@auriniapharma.com 501-590-3690

> Anna Pavlova-Wolf

Medical Science Liaison apavlova@auriniapharma.com 903-315-0080

#### **Bristol Myers Squibb Contacts:**

> Karen Smith

Karen.smith1@bms.com 918-857-3807

> Karl Friday

Karl.Friday@bms.com 918-671-9400

#### **GlaxoSmithKline Contacts:**

Margie Monroe

405.209.4601 margie.l.monroe@gsk.com

#### Janssen/Johnson & Johnson

> Jerome Reed

jreed29@its.jnj.com 513-309-8972

> Scott Jackson

Sjacks19@its.jnj.com 619-988-0482

## Exhibitors' Virtual Zoom Marketplace

7:30am-8:30am | 11:am (morning break) | 12:15 (lunch)



https://gabs.zoom.us/j/84385988531?pwd=R1M2WU1mOTdlbnFpRlRnMmhJeWJZQT09

Meeting ID: 843 8598 8531

Passcode: 271233



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Meeting ID: 875 1622 2263

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Meeting ID: 886 2139 6370

Passcode: 271233



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Meeting ID: 818 5112 5098

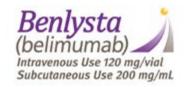
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Meeting ID: 846 0586 9689

Passcode: 271233



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Meeting ID: 826 9635 4039

Passcode: 320318



https://gabs.zoom.us/j/84385988531?pwd=R1M2WU1mOTdlbnFpRIRnMmhJeWJZQT09

Meeting ID: 843 8598 8531

Passcode: 271233

# Connect Perplexing Symptoms With Low Alkaline Phosphatase

The diagnosis may point to hypophosphatasia<sup>1</sup>

Ordered Items

Comp. Metabolic Panel (14): Venipuncture

TESTS	RESULT	FLAG
Comp. Metabolic Panel (14)		
Glucose, serum	75	
BUN	10	
Creatinine, serum	1.0	
Sodium, serum	140	
Potassium, serum	3.9	
Chloride, serum	102	
Carbon dioxide, total	26	
Calcium, serum	10.4	
Albumin, serum	3.7	
Total protein	6.5	
Bilirubin, total	0.7	
AST (SGOT)	15	
ALT (SGPT)	17	
Alkaline phosphatase	31	Low

Persistently low alkaline phosphatase (alk phos) levels, <40 U/L, can be the key to identifying metabolic bone disease in hard-to-diagnose patients.  $^{1.3\ast}$ 

Patients with hypophosphatasia (HPP) may experience one or more key symptoms, including joint, bone, or muscle pain, and muscle weakness, along with mobility issues, frequent fractures, and poor dentition. When patients present with one or more key symptoms, or report them in questions on background history, the next step is to check for low alk phos on their comprehensive metabolic panel.

Patient image and patient lab values are hypothetical. ALP reference ranges vary based upon lab. Refer to your lab for the appropriate reference intervals.

\*Limitations: An alk phos level of below 40 U/L is not conclusive for a diagnosis of HPP. Patient should be evaluated for other symptoms of HPP and differential diagnoses should be ruled out.

References: I. Bilanchi M.L. Bishop N.J. Guañabens N. et al. Osteoporns Int. 2020;3[8]:1445-1460. 2. Adeli K. Higgins V. Nieuwesteeg M. et al. Clin Chem. 2015;6[8]:1049-1062. 3. Schumann G. Klauke R. Canalias F, et al. Clin Chem Lob Med. 2011;49[9]:1439-1446.

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<sup>a</sup>Complete renal response was achieved in 40.8% of patients with LUPKYNIS and 22.5% with control. Proteinuria reductions (UPCR ≤0.5 mg/mg) were achieved at a median time of 169 days with LUPKYNIS vs 372 days with control.¹

bComplete renal response was defined as a confirmed UPCR of ≤0.5 mg/mg; eGFR ≥60 mL/min/1.73 m² or no confirmed decrease from baseline in eGFR of >20% or no treatment- or disease-related eGFR-associated event at time of assessment; presence of sustained, low-dose steroids (≤10 mg prednisone from Weeks 44-52); and no administration of rescue medications. Proteinuria reduction was based on time to UPCR of ≤0.5 mg/mg.¹



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Questions? Call 1-833-AURINIA (1-833-287-4642) 8AM to 8PM ET, fax to 1-833-213-1001, or email support@AuriniaAlliance.com

eGFR=estimated glomerular filtration rate; MMF=mycophenolate mofetil; standard of care=MMF + steroids; UPCR=urine protein/creatinine ratio.

#### **Indications**

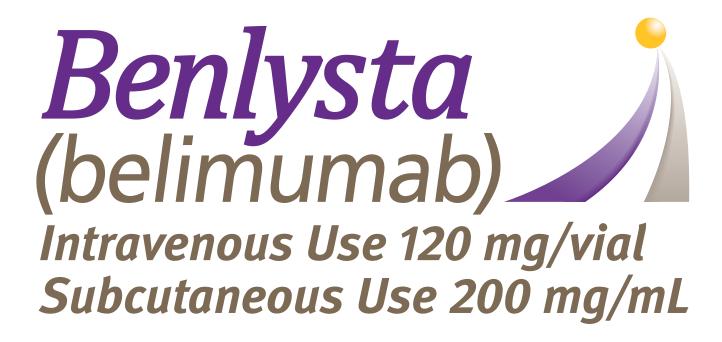
LUPKYNIS is indicated in combination with a background immunosuppressive therapy regimen for the treatment of adult patients with active lupus nephritis (LN). Limitations of Use: Safety and efficacy of LUPKYNIS have not been established in combination with cyclophosphamide. Use of LUPKYNIS is not recommended in this situation.

#### **Important Safety Information**

BOXED WARNINGS: MALIGNANCIES AND SERIOUS INFECTIONS Increased risk for developing malignancies and serious infections with LUPKYNIS or other immunosuppressants that may lead to hospitalization or death.

Please see additional <u>Important Safety Information</u> and <u>Prescribing Information</u> including Boxed Warning and Medication Guide for LUPKYNIS.







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