Bimekizumab Efficacy in High-Impact Areas: Pooled

2-Year Analysis in Scalp, Nail, and Palmoplantar Psoriasis
from Phase 3/3b Randomized Controlled Trials

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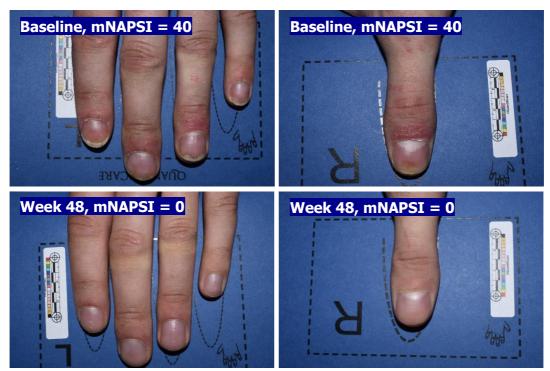
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Fig S1. Bimekizumab nail treatment example over 48 weeks from BE RADIANT



Previously presented at EADV 2021. Photos show the left-hand fingers and right thumb from one patient treated with bimekizumab (BKZ) every 4 weeks (Q4W)/every 8 weeks (Q8W) in BE RADIANT and included in this analysis, with the total modified Nail Psoriasis Severity Index (mNAPSI) shown at baseline and Week 48. BKZ: bimekizumab; mNAPSI: modified Nail Psoriasis Severity Index; Q4W: every 4 weeks; Q8W: every 8 weeks.

Table S1. Mean percent change from baseline in scalp, nail, or palmoplantar psoriasis over 2 years (OC)

| | Scalp IGA ≥3 (N=821) | | mNAPSI >10 (N=377) | | Palmoplantar IGA ≥3 (N=193) | |
|--------------------------------------|-------------------------|---------------------------|-----------------------|---------------------------|--------------------------------|--------------------------|
| OC; mean % change (SE) | BKZ total (N=821) | BKZ Q4W/Q8W (n=277) | BKZ total (N=377) | BKZ Q4W/Q8W (n=129) | BKZ total (N=193) | BKZ Q4W/Q8W (n=52) |
| Week 16 | -92.2 (0.7) | -93.5 (1.1) | -64.5 (2.0) | -71.1 (2.7) | -92.6 (1.5) | -97.0 (1.3) |
| Year 1 (OLE Week 0 ^a) | -94.1 (0.7) | -96.3 (0.8) | -91.3 (1.1) | -93.2 (1.4) | -95.3 (1.2) | -97.6 (1.5) |
| OLE Week 24 | -93.4 (0.7) | -95.3 (1.0) | -93.4 (0.9) | -94.8 (1.1) | -96.7 (1.1) | -95.3 (2.4) |
| Year 2 (OLE Week 48) | -93.4 (0.7) | -94.6 (1.2) | -91.5 (1.3) | -92.5 (1.7) | -96.2 (1.1) | -97.2 (1.4) |

Scalp Investigator's Global Assessment (IGA) data are for patients with baseline scalp IGA ≥3, modified Nail Psoriasis Severity Index (mNAPSI) for patients with baseline mNAPSI >10, and palmoplantar IGA for patients with baseline palmoplantar IGA ≥3. Data are presented for patients with plaque psoriasis initially randomized to bimekizumab (BKZ) who later entered the open-label extension (OLE). Observed case (OC) was used for missing data. BKZ total consists of patients randomized to receive BKZ 320 mg every 4 weeks (Q4W) to Week 16, and who received either BKZ Q4W or every 8 weeks (Q8W) to the end of the first year (Week 48/52/56), and entered the OLE. BKZ Q4W/Q8W consists of patients randomized to BKZ 320 mg Q4W to Week 16, who received BKZ Q8W throughout the maintenance period and on OLE entry; there are no BE VIVID patients in this treatment arm. [a] OLE Week 0 corresponds to Week 48 for BE SURE, BE READY, and BE RADIANT, and Week 52 for BE VIVID. BKZ: bimekizumab; IGA: Investigator's Global Assessment; mNAPSI: modified Nail Psoriasis Severity Index; OC: observed case; OLE: open-label extension; Q4W: every 4 weeks; Q8W: every 8 weeks; SE: standard error.

Table S2. Complete clearance in scalp, nail, or palmoplantar psoriasis over 2 years (NRI)

| | Scalp IGA ≥3 (N=821) | | mNAPSI >10 (N=377) | | Palmoplantar IGA ≥3 (N=193) | |
|--|-------------------------|---------------------------|-----------------------|---------------------------|--------------------------------|--------------------------|
| Achievement of scalp IGA 0, mNAPSI 0, or palmoplantar IGA 0, n (%) | BKZ total (N=821) | BKZ Q4W/Q8W (n=277) | BKZ total (N=377) | BKZ Q4W/Q8W (n=129) | BKZ total (N=193) | BKZ Q4W/Q8W (n=52) |
| Week 16 | 681 (82.9) | 239 (86.3) | 85 (22.5) | 39 (30.2) | 167 (86.5) | 47 (90.4) |
| Year 1 (OLE Week 0 ^a) | 696 (84.8) | 245 (88.4) | 237 (62.9) | 89 (69.0) | 167 (86.5) | 49 (94.2) |
| OLE Week 24 | 630 (76.7) | 220 (79.4) | 243 (64.5) | 89 (69.0) | 140 (72.5) | 35 (67.3) |
| Year 2 (OLE Week 48) | 643 (78.3) | 230 (83.0) | 246 (65.3) | 87 (67.4) | 151 (78.2) | 43 (82.7) |

Scalp Investigator's Global Assessment (IGA) 0 data are presented for patients with baseline scalp IGA ≥3, modified Nail Psoriasis Severity Index (mNAPSI) 0 for patients with baseline mNAPSI >10, and palmoplantar IGA 0 for patients with baseline palmoplantar IGA ≥3. Data are presented for patients with plaque psoriasis initially randomized to BKZ who later entered the open-label extension (OLE). Non-responder imputation (NRI) was used for missing data. BKZ total consists of patients randomized to receive BKZ 320 mg every 4 weeks (Q4W) to Week 16, and who received either BKZ Q4W or every 8 weeks (Q8W) to the end of the first year (Week 48/52/56), and entered the OLE. BKZ Q4W/Q8W consists of patients randomized to BKZ 320 mg Q4W to Week 16, who received BKZ Q8W throughout the maintenance period and on OLE entry; there are no BE VIVID patients in this treatment arm. [a] OLE Week 0 corresponds to Week 48 for BE SURE, BE READY, and BE RADIANT, and Week 52 for BE VIVID. BKZ: bimekizumab; IGA: Investigator's Global Assessment; mNAPSI: modified Nail Psoriasis Severity Index; NRI: non-responder imputation; OC: observed case; OLE: open-label extension; Q4W: every 4 weeks; Q8W: every 8 weeks.