

SCEMBLIX - The unmet need in CML - HCP

Scemblix® (asciminib) prescribing information Glivec® (imatinib) prescribing information

Image



Image



SCEMBLIX®  $\blacktriangledown$  (asciminib) is indicated for the treatment of adult patients with Philadelphia chromosome-positive chronic myeloid leukaemia (Ph + CML) in chronic phase (CP), previously treated with two or more tyrosine kinase inhibitors, and without a known T315I mutation.  $^1$ 

# The unmet need in chronic myeloid leukaemia

Tyrosine kinase inhibitors (TKIs) have improved CML prognosis, yet survival remains poor in patients on  $\geq$ 3rd line therapy.<sup>2,3</sup>

Many patients on TKIs face treatment failure or discontinuation due to intolerance, with up

to half of patients discontinuing 1st line imatinib within 5 years.3

During 2nd-line treatment:<sup>3</sup>



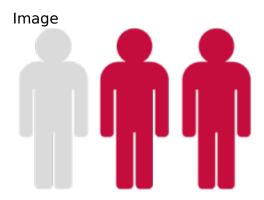
of patients fail to achieve major molecular response (MMR)  $(n=1279)^{*3}$ 



of patients fail to achieve complete cytogenetic response (CCyR) within 2 years  $(n=958)^{*3}$ 

\*Results for MMR and CCyR rates are taken from multiple studies including different 2nd line medications. This is a meta-analysis.<sup>3</sup>

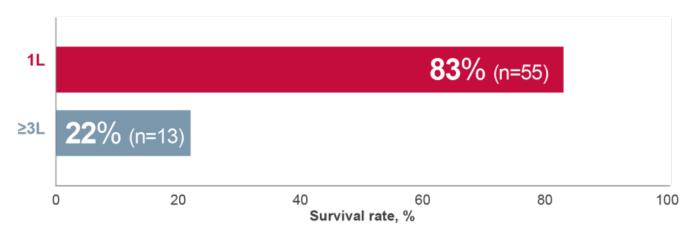
In CML patients, a low 8-year overall survival was associated with being on  $\geq$ 3rd line treatment.<sup>†2</sup>



In approximately 2/3 of those who had at least 1 TKI switch it was due to  $resistance^{t4}$  (n=73/113)

## 8-year overall survival, N=902

**Image** 



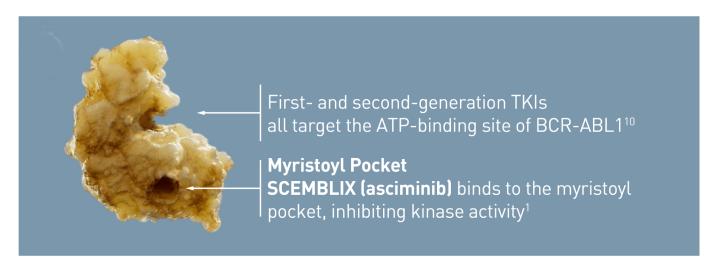
Adapted from Bosi GR, et al. 2019.2



Optimal monitoring is important to assess treatment benefits and inform the decision to switch.<sup>5,6</sup> When a TKI fails, timely switching can limit the progression of disease<sup>6</sup>

A 2nd generation TKI may have limited benefit in a ≥3L setting, after the failure of another 2nd generation TKI and 1st generation TKI prior.<sup>3</sup> Patients in ≥3L treatment may need a different mechanism of action to optimise outcomes.<sup>2,7-9</sup>

**Image** 



"You can end up running out of options for patients who experience side effects. Then you need to consider transplant after 4th line or chemotherapy options"

### Adapted from haematologist quote<sup>11</sup>

"In a year I had gone from taking one drug and living a normal life to having zero options"

Adapted from 5L CML patient quote<sup>12</sup>

# SCEMBLIX is the first and only STAMP inhibitor, specifically targeting the ABL1 myristoyl pocket<sup>10,13,14</sup>

### Discover the MOA

The ASCEMBL trial did not restrict to Ph+ patients with CP-CML. SCEMBLIX is indicated in adults with Ph+ CP-CML previously treated with two or more tyrosine kinase inhibitors and without a known T315I mutation.<sup>1,13</sup>

<sup>†</sup>Data from a retrospective, non-interventional study conducted at 21 UK NHS secondary and tertiary care centres on 257 patients with CML.<sup>4</sup>

1L, first line; 3L, third line; ATP, adenosine triphosphate; BCR-ABL, breakpoint cluster region and Abelson murine leukaemia viral oncogene homologue; CCyR, complete cytogenetic response; CI, confidence interval; CML, chronic myeloid leukaemia; CP, chronic phase; MCyR, major cytogenetic response; MMR, major molecular response; MOA, mechanism of action; NHS, National Health Service; Ph+, Philadelphia chromosome positive; STAMP, specifically targeting the ABL1 myristoyl pocket; TKI, tyrosine kinase inhibitor.

For further information, please refer to the **Summary of Product Characteristics**.

#### References

1. SCEMBLIX (asciminib) Summary of Product Characteristics.

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