CD157 signaling promotes survival of acute myeloid leukemia cells through regulation of Mcl-1

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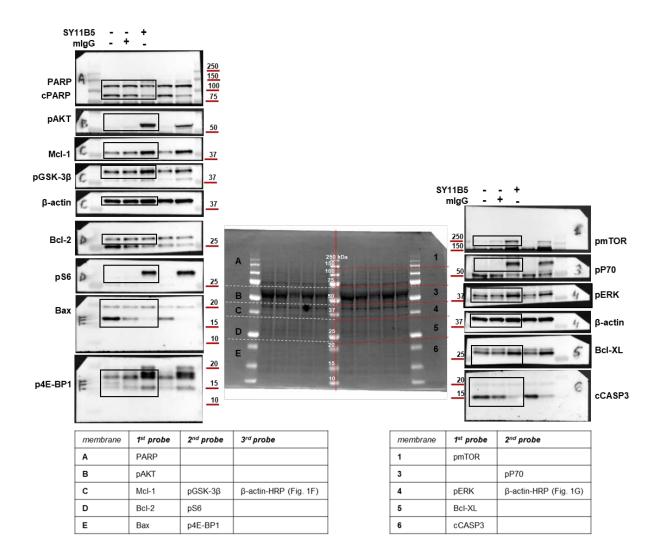
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Uncropped blot images

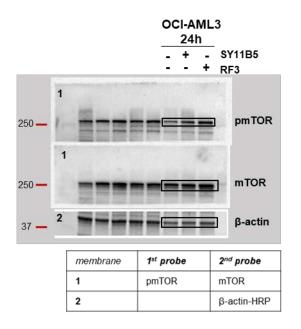
Supplementary Fig. S8

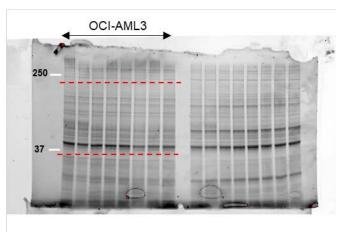
Figure 1F/1G

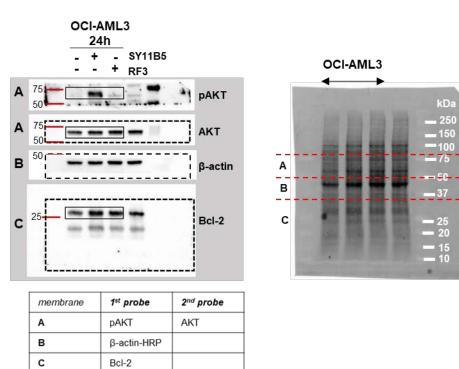


Uncropped blot from western blot analysis presented in Figure 1F-G. Panel F and G were generated from the same blot using the same samples. Dotted lines in full-length blot indicate the excision lines. Each strip was separately probed with the indicated antibodies. After hybridization with the first probe, the membrane was stripped and re-hybridized with the second probe according to the table under each blot. The rectangles show the cropped region of the blots included in Fig. 1F-G.

Figure 2B

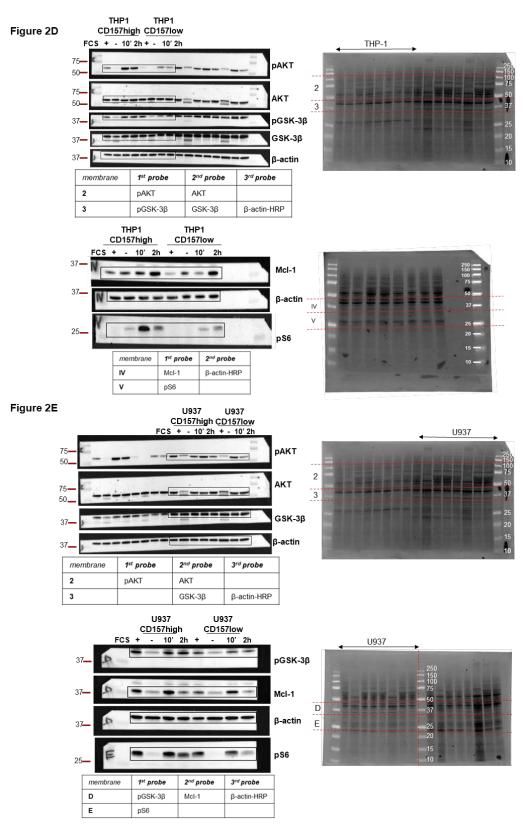






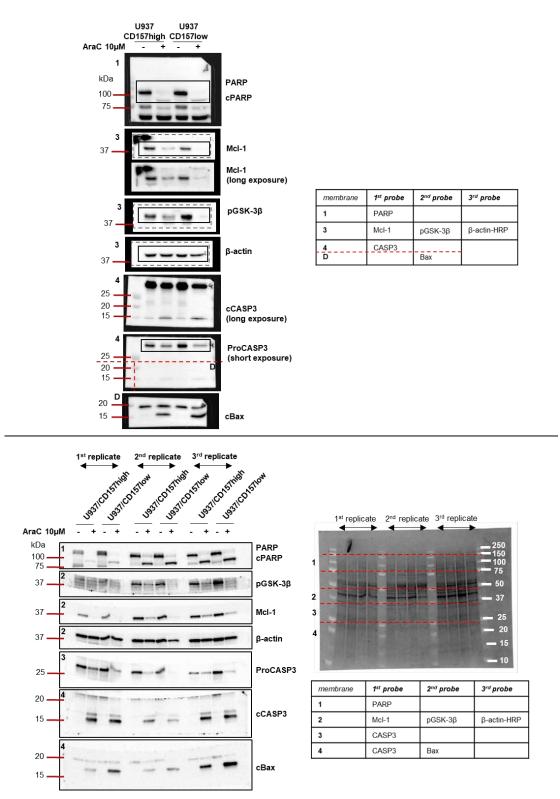
Uncropped gel (top) and blot (bottom) from western blot analysis presented in Figure 2B. Two separate blots were generated using the same samples. Dotted lines in uncropped images indicate the excision lines. Each strip was separately probed with the indicated antibodies. After hybridization with first probe, the membrane was stripped and re-hybridized with the second probe according to the table under each blot. The rectangles show the cropped region of the blots included in the figure. Black dotted lines indicate membrane periphery.

Supplementary Fig. S9 (continued)



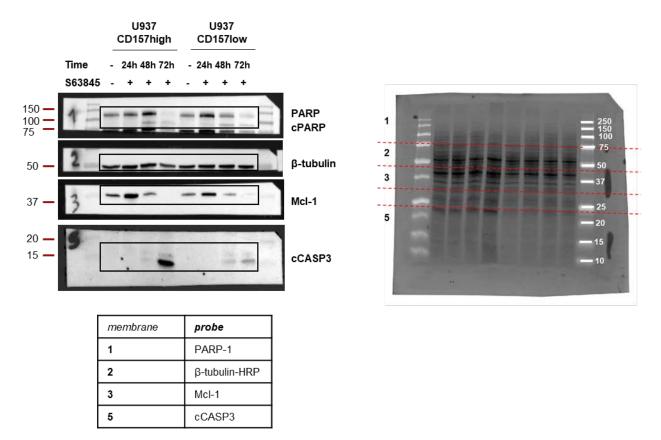
Uncropped blots from western blot analysis presented in Figure 2D-E. For each cell line two separate blots were generated using the same samples. Dotted red lines in full-length blot indicate the excision lines. Each strip was separately probed with the indicated antibodies. After hybridization with first probe, the membrane was stripped and re-hybridized with the second probe according to the table under each blot. The rectangles show the cropped region of the blots included in the figure.

Figure 3D



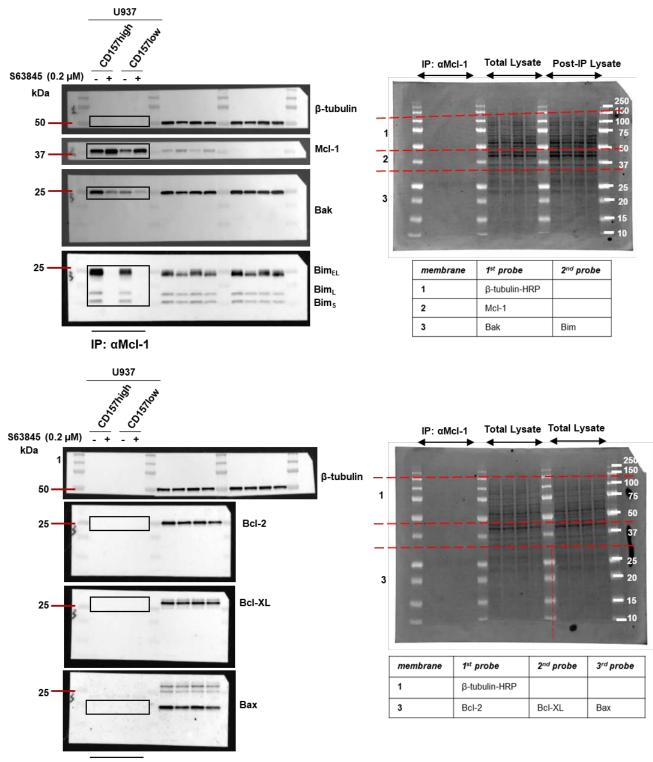
Uncropped blot presented in Figure 3D. Upper blot was cut prior to hybridization and each strip was separately probed with the indicated antibodies. After hybridization with first probe, the membrane was stripped and re-hybridized with the second probe according to each table. The rectangles show the cropped region of the blots included in Fig. 3D. Multiple exposures are shown for blots that were taken with a higher contrast or do not have background sufficiently dark to show the membrane periphery. Grey dotted lines indicate membrane periphery. Bottom blot show three independent experiments run in parallel and used for quantification of protein levels by densitometry analysis reported in Fig. 3D (left panel).

Figure 4B



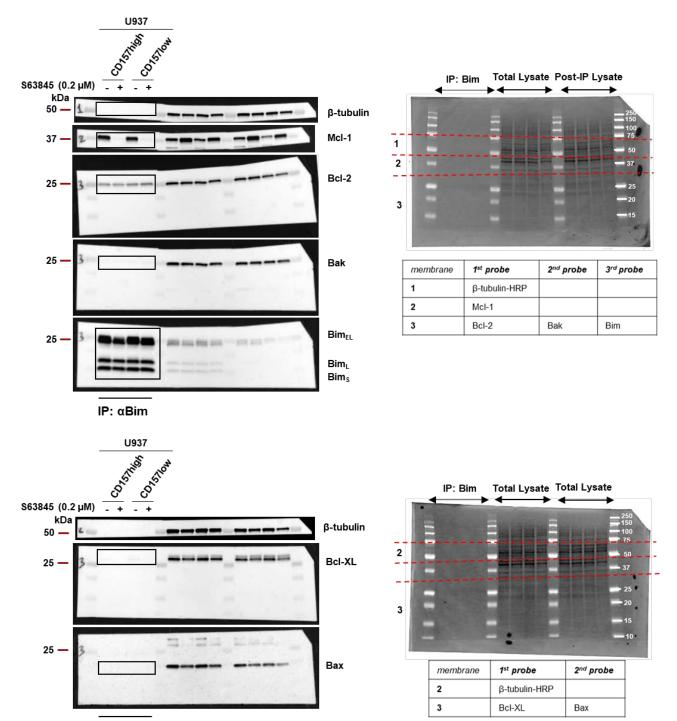
Uncropped version of blot presented in Figure 4B. The rectangles show the cropped region of the blots included in the figure. Dotted red lines in full-length blot indicate the excision lines. Each strip was separately probed with the indicated antibody.

Figure 4D



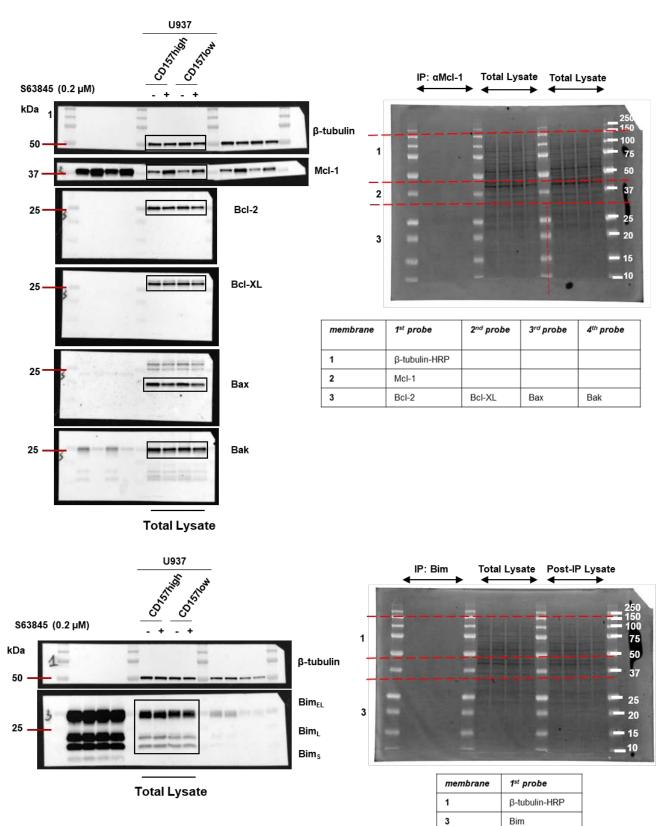
IP: αMcI-1

Figure 4D



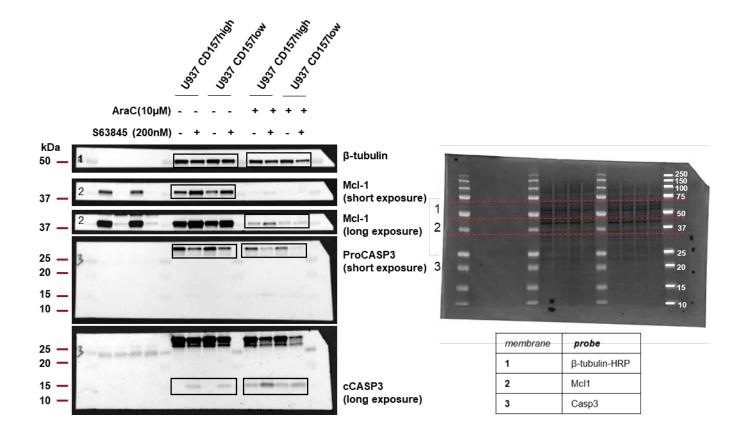
IP: αBim

Figure 4D



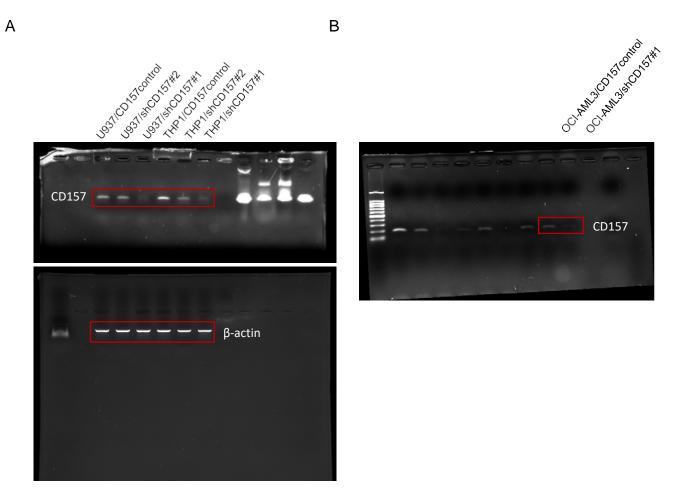
Uncropped blot from the western blot analysis presented in Figure 4D. All blots were generated using the same samples. Dotted red lines in uncropped images indicate the excision lines. Each strip was separately probed with the indicated antibodies. After hybridization with first probe, the membrane was stripped and re-hybridized with the second probe according to the table under each blot. The rectangles show the cropped region of the blots included in Fig. 4D.

Figure 5C



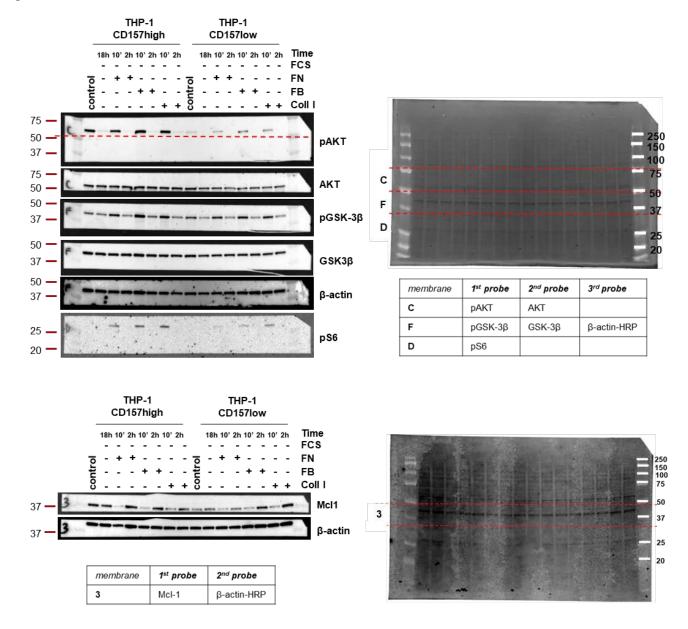
Uncropped blot from the western blot analysis presented in Figure 5C. Dotted lines in uncropped images indicate the excision lines. Each strip was separately probed with the indicated antibody. Multiple exposures are shown for blots that were taken with a higher contrast The rectangles show the cropped region of the blots included in Fig. 5C.

Figure S3



Uncropped gels from PCR analysis performed in Figure S3. The rectangles show the cropped region of the gels included in Fig. S3.

Figure S5



Uncropped blots from western blot analysis presented in Figure S5. Two separate blots were generated using the same samples. Dotted red lines in full-length blots indicate the excision lines. Each strip was separately probed with the indicated antibodies. After hybridization with first probe, the membrane was stripped and re-hybridized with the second probe according to the table under each blot.