

# Human SLC39A14 (ZIP-14) knockout HEK-293T cell line ab266126

3 Images

### Overview

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<b>Product name</b>	Human SLC39A14 (ZIP-14) knockout HEK-293T cell line
<b>Parental Cell Line</b>	HEK293T
<b>Organism</b>	Human
<b>Mutation description</b>	Knockout achieved by using CRISPR/Cas9, Homozygous: 1 bp insertion in exon 2
<b>Passage number</b>	<20
<b>Knockout validation</b>	Sanger Sequencing, Western Blot (WB)
<b>Tested applications</b>	<b>Suitable for:</b> WB, Sanger Sequencing
<b>Biosafety level</b>	2
<b>General notes</b>	<p><b>Recommended control:</b> Human wild-type HEK293T cell line (<a href="#">ab255449</a>). Please note a wild-type cell line is not automatically included with a knockout cell line order, if required please add recommended wild-type cell line at no additional cost using the code WILDTYPE-TMTK1.</p> <p><b>Cryopreservation cell medium:</b> Cell Freezing Medium-DMSO Serum free media, contains 8.7% DMSO in MEM supplemented with methyl cellulose.</p> <p><b>Culture medium:</b> DMEM (High Glucose) + 10% FBS</p> <p><b>Initial handling guidelines:</b> Upon arrival, the vial should be stored in liquid nitrogen vapor phase and not at -80°C. Storage at -80°C may result in loss of viability.</p> <ol style="list-style-type: none"><li>1. Thaw the vial in 37°C water bath for approximately 1-2 minutes.</li><li>2. Transfer the cell suspension (0.8 mL) to a 15 mL/50 mL conical sterile polypropylene centrifuge tube containing 8.4 mL pre-warmed culture medium, wash vial with an additional 0.8 mL culture medium (total volume 10 mL) to collect remaining cells, and centrifuge at 201 x g (rcf) for 5 minutes at room temperature. 10 mL represents minimum recommended dilution. 20 mL represents maximum recommended dilution.</li><li>3. Resuspend the cell pellet in 5 mL pre-warmed culture medium and count using a haemocytometer or alternative cell counting method. Based on cell count, seed cells in an appropriate cell culture flask at a density of <math>2 \times 10^4</math> cells/cm<sup>2</sup>. Seeding density is given as a guide only and should be scaled to align with individual lab schedules.</li><li>4. Incubate the culture at 37°C incubator with 5% CO<sub>2</sub>. Cultures should be monitored daily.</li></ol> <p><b>Subculture guidelines:</b></p> <p>All seeding densities should be based on cell counts gained by established methods. A guide seeding density of <math>2 \times 10^4</math> cells/cm<sup>2</sup> is recommended.</p>

A partial media change 24 hours prior to subculture may be helpful to encourage growth, if required.

Cells should be passaged when they have achieved 80-90% confluence.

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We will provide viable cells that proliferate on revival.

## Properties

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<b>Number of cells</b>	1 x 10 <sup>6</sup> cells/vial, 1 mL
<b>Adherent /Suspension</b>	Adherent
<b>Tissue</b>	Kidney
<b>Cell type</b>	epithelial
<b>STR Analysis</b>	Amelogenin X D5S818: 8, 9 D13S317: 12, 14 D7S820: 11 D16S539: 9, 13 vWA: 16, 19 TH01: 7, 9.3 TPOX: 11 CSF1PO: 11, 12
<b>Mycoplasma free</b>	Yes
<b>Storage instructions</b>	Shipped on Dry Ice. Store in liquid nitrogen.
<b>Storage buffer</b>	Constituents: 8.7% Dimethylsulfoxide, 2% Cellulose, methyl ether

## Target

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<b>Function</b>	May be able to transport iron (By similarity). Acts as a zinc-influx transporter.
<b>Tissue specificity</b>	Ubiquitously expressed, with increased expression in liver, pancreas, fetal liver, thyroid gland, left and right ventricle, right atrium and fetal heart. Weakly expressed in spleen, thymus, and peripheral blood leukocytes.
<b>Sequence similarities</b>	Belongs to the ZIP transporter (TC 2.A.5) family.
<b>Cellular localization</b>	Cell membrane. Cell projection > lamellipodium. Localized to the plasma membrane and also found colocalized with F-actin concentrated on lamellipodiae.

## Applications

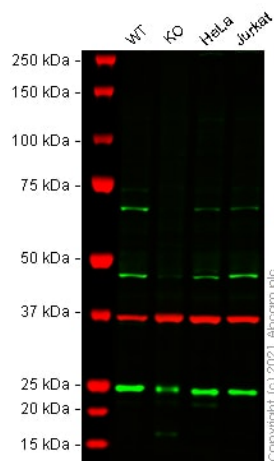
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**The Abpromise guarantee** Our [Abpromise guarantee](#) covers the use of ab266126 in the following tested applications. The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
<b>WB</b>		Use at an assay dependent concentration. Predicted molecular weight: 54 kDa.
<b>Sanger Sequencing</b>		Use at an assay dependent concentration.

## Images

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Western blot - Human SLC39A14 (ZIP-14) knockout HEK-293T cell line (ab266126)

**All lanes :** Anti-SLC39A14/ZIP-14 antibody (**ab106568**)

**Lane 1 :** Wild-type HEK-293T cell lysate

**Lane 2 :** SLC39A14 knockout HEK-293T cell lysate

**Lane 3 :** HeLa cell lysate

**Lane 4 :** Jurkat cell lysate

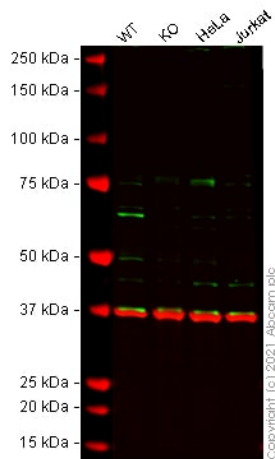
Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

**Predicted band size:** 54 kDa

**Observed band size:** 68 kDa

False colour image of Western blot: Anti-SLC39A14/ZIP-14 antibody staining at 1 µg/ml, shown in green; Mouse anti-GAPDH antibody [6C5] (**ab8245**) loading control staining at 1/20000 dilution, shown in red. In Western blot, **ab106568** was shown to bind specifically to SLC39A14/ZIP-14. A band was observed at 68 kDa in wild-type HEK-293T cell lysates with no signal observed at this size in SLC39A14 knockout cell line ab266126 (knockout cell lysate **ab258683**). To generate this image, wild-type and SLC39A14 knockout HEK-293T cell lysates were analysed. First, samples were run on an SDS-PAGE gel then transferred onto a nitrocellulose membrane. Membranes were blocked in 3 % milk in TBS-0.1 % Tween<sup>®</sup>20 (TBS-T) before incubation with primary antibodies overnight at 4 °C. Blots were washed four times in TBS-T, incubated with secondary antibodies for 1 h at room temperature, washed again four times then imaged. Secondary antibodies used were Goat anti-Rabbit IgG H&L (IRDye<sup>®</sup> 800CW) preabsorbed (**ab216773**) and Goat anti-Mouse IgG H&L (IRDye<sup>®</sup> 680RD) preabsorbed (**ab216776**) at 1/20000 dilution.



Western blot - Human SLC39A14 (ZIP-14) knockout HEK-293T cell line (ab266126)

**All lanes** : Anti-SLC39A14/ZIP-14 antibody ([ab191199](#)) at 1/2000 dilution

**Lane 1** : Wild-type HEK-293T cell lysate

**Lane 2** : SLC39A14 knockout HEK-293T cell lysate

**Lane 3** : HeLa cell lysate

**Lane 4** : Jurkat cell lysate

Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

**Predicted band size:** 54 kDa

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False colour image of Western blot: Anti-SLC39A14/ZIP-14 antibody staining at 1/2000 dilution, shown in green; Mouse anti-GAPDH antibody [6C5] ([ab8245](#)) loading control staining at 1/20000 dilution, shown in red. In Western blot, [ab191199](#) was shown to bind specifically to SLC39A14/ZIP-14. A band was observed at 68 kDa in wild-type HEK-293T cell lysates with no signal observed at this size in SLC39A14 knockout cell line ab266126 (knockout cell lysate [ab258683](#)). To generate this image, wild-type and SLC39A14 knockout HEK-293T cell lysates were analysed. First, samples were run on an SDS-PAGE gel then transferred onto a nitrocellulose membrane. Membranes were blocked in 3 % milk in TBS-0.1 % Tween<sup>®</sup>20 (TBS-T) before incubation with primary antibodies overnight at 4 °C. Blots were washed four times in TBS-T, incubated with secondary antibodies for 1 h at room temperature, washed again four times then imaged. Secondary antibodies used were Goat anti-Rabbit IgG H&L (IRDye<sup>®</sup> 800CW) preabsorbed ([ab216773](#)) and Goat anti-Mouse IgG H&L (IRDye<sup>®</sup> 680RD) preabsorbed ([ab216776](#)) at 1/20000 dilution.

Mut	GAGCCTCAGGGGTGGTTCTCCATAAGCCAAAGCAGGGTCAGCAGGAGGCAGCTCTGGAAG
WT	GAGCCTCAGGGGTGGTTCTCCATAAGCCAA GCAGGGTCAGCAGGAGGCAGCTCTGGAAG
Sanger Sequencing - Human SLC39A14 knockout	
HEK293T cell line (ab266126)	

Homozygous: 1 bp insertion in exon 2

**Please note:** All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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