Working at home is a privilege

The chief executive officer of Yahoo stirred up a bit of controversy this week after laying down the law about working from home.

Marissa Meyer was appointed CEO in July and has the challenging assignment of making sure Yahoo

doesn't go
the way of
Magellan or
Infoseek or
AltaVista.
(For younger
readers
these were
search
engines in
prehistoric
Internet



Rebecca Regnier

times that have gone the way of the pterodactyl.)

The boss of Yahoo pulled the plug on "working from home" or telecommuting arrangements. By June all Yahoo employees who work from home need to get to the office.

This notion that Yahoo employees should be required to go to the office has been criticized as "crusty," "antiquated," "janked up." Work-athome proponents say flexibility promotes a more stress-free life, that it's good for families, the environment and pajama sales! Sorry, that last thing is a stereotype typically associated with working at home. You also can wear yoga pants.

The internal memo, leaked to a Web site called All Things D, explained that being in the same place with your co-workers promotes collaboration, innovation and showering! Sorry that last thing is a stereotype associated with working at home. Going into the office is no guarantee that everyone has showered.

First off, where was I when they were handing out work-at-home jobs? I've got skills, I'm internally

motivated, I'm busy with family crap, and I have a wireless modem. Pick ME.

Before I go any further, I understand that working at home is a thing and that some people do it well. I tip my hat to them. I could never find a company willing to let me do the same. I am jealous.

That said. Shut the heck up. So many creative, hardworking and qualified people have no work at all. Say what you will about CEOs but Ms. Meyer's plan to put the best minds in one place to give Yahoo a fighting chance seems pretty smart to me. No one will be working at Yahoo from home or otherwise if she fails.

This one time I had a baby, the next day my husband went back to work. The next week I did. Why, because the luxury of working at home wasn't an option and make no mistake, it's a luxury. One that, right now, Yahoo can't afford.

And, yes, I wrote this column from home, I have on yoga pants and I'm supervising one of my kid's attempts at making lasagna. I wrote it after an eight-hour day at the office, the one with the health benefits. Am I asking for pity? No, I'm saying suck it up Yahoo work-athomers — you have a job.

Finally just because one company institutes a moratorium doesn't signal the end of working at home. Technology means that working at home is an increasingly more viable option. Working at home isn't going away.

As the controversy boiled up, Yahoo issued this statement: "We don't discuss internal matters. This isn't a broad industry view on working from home — this is about what is right for Yahoo!, right now."

Right on.

Making connections

Traveling box links families of children with rare disease

BY PAULA WETHINGTON paula@monroenews.com

In a few days, an Ohio family will get a traveling care package from Monroe.

It will include a little girl's play dress, a bright green tie for a boy, cards, photos and other messages meant for families whose children have been diagnosed with a spontaneous genetic disorder called cyclin-dependent kinase-like 5, which is usually abbreviated as CDKL5.

Last Friday, the package was delivered to Jakob Cooper, 9, a student at Monroe County Intermediate School District, and his family: parents Robert and Tammy Cooper, brother Jayce, 7, and sisters Sarah, 19, and Stefanie, 23.

"I got kind of emotional when I opened the box," Mrs. Cooper said Tuesday.

The project, which started in summer, 2011, is called Little Green Dress, and is coordinated by International Foundation for CDKL5 Research. Each family is invited to enclose a picture of their daughter wearing the green dress and accompanying wreath or their son wearing the green tie, sign a traveling journal, pick up a gift and supply a small gift to the next child such as a stuffed animal.

The Coopers are the 21st family on the list. The box so far has been in the United Kingdom, Denmark, Illinois and Minnesota. The next family lives in Ohio, another that is on the list lives in Japan.

Mrs. Cooper said one of the things that amazed her, upon inspection of the pictures and stories, was "how different all



Evening News photo by KIM BREN

Tamara Cooper and her son, Jakob Cooper, 9, are the most recent recipients of a package containing photographs and journal entries from families around the world with children affected by CDKL5, a rare disease that was only discovered early in the 21st century.

these kids are!"

Jakob was about a month old when his mother started to become concerned about his situation.

"He was my third child, so we knew something was wrong," she said.

Over time, they noticed his conditions include uncontrolled seizures and severe visual impairment. It took a year to get the first diagnosis: mitochondrial disease complex IV deficiency.

But that was rewritten after Mrs. Cooper realized her son was not following the expected path of children with that condition.

About a year and a half



The idea of the travelling box, which includes a journal, green necktie and green dress representing the disease, started in the United Kingdom and has made its way through Canada and into the United States.

ago, a doctor re-evaluated Jakob's situation and it has now been pinpointed to CDKL5. This doesn't change his medications or care plan.

But it does provide the family with some answers.

One of the reasons that

RARE DISEASE DAY

Today is the Sixth International Rare Disease Day, with organizations in more than 60 countries campaigning to bring attention to conditions that affect only a few people.

The goal is to encourage and support coordination and cooperation among research teams whose expertise may be scattered across multiple countries.

For more information, visit www.cdkl5.com and www.rarediseaseday.org.

condition was not picked up on at first is that it is extremely rare. Even a year ago, there were only about 200 children known to have it. Shortly after Mrs. Cooper posted on her Facebook page that was the diagnosis, she got a message from a mom who is involved with the foundation and started to make friends online with others in the situation.

The condition also is more likely to happen in girls than in boys, although Mrs. Cooper said boys who are affected have far more serious disabilities.

As she looked through the pictures of children whose families are participating in the Little Green Dress project, she pointed out that the boys were more likely to be seated in wheelchairs than the girls.

And she felt connections to those families, regardless of where they live.

"Thank you for letting us be part of this terrific journey of this beautiful green dress," she wrote in the traveling journal. "Hugs and kisses, Tammy Cooper."

STATE LEGISLATURE

Lawmaker proposes lead testing for children

LANSING (AP) — One Michigan lawmaker wants young children in the state to be screened for lead poisoning.

Democratic Rep. Thomas Stallworth of Detroit introduced a bill Wednesday that would require primary health-care professionals to screen all 1- to 2-yearold patients for lead poisoning.

If a child meets certain criteria, he or she would also be given a blood lead diagnostic test.

Parents who object because of religious beliefs can opt out.

The Center for Disease Control and Prevention says lead-based paint and dust that has been contaminated with lead are the primary sources of lead poisoning among children.

It says children are particularly susceptible to poisoning because they are more likely to touch objects contaminated with lead and then put their hands in their mouths.



